

Single Registration Number (SRN): N/A



Urgent Product Correction

Immediate Action Required

Date Issued

September 29, 2021

Product

Product Description	List Number	Serial Number	US UDI	EU UDI
ARCHITECT i1000SR	1L86 1L87	See Attachment A	N/A	N/A
ARCHITECT i2000SR	3M74			
ARCHITECT i2000	1G17 8C89			
ARCHITECT c4000	2P24 1P86 1R24 1R25			
ARCHITECT c8000	1G06			
ARCHITECT c16000	3L77			

Explanation

Abbott has identified three potential performance issues for the ARCHITECT Software version 9.41 and earlier. Abbott is releasing ARCHITECT Software versions 9.45 and 9.50 to correct these issues (see details in **Appendix A**).

1. When an Error Code 3382, 'Unable to process test, internal wash pressure (x) error (y) pipettor' occurs, the ARCHITECT c4000 and the ARCHITECT c16000 are incorrectly placed into a 'Scheduled Pause' status rather than a 'Stopped' status. As a result, the processing module continues to process tests after the hardware error is detected. This may cause incorrect results to be generated.
2. When configuring a Calibrator Sample Volume on the Configure Assays screen, if the user selects multiple assays, the calibrator sample volume from one assay may be carried into the calibrator sample volume for another assay. Incorrect calibrator sample volumes have the potential to generate incorrect calibration curves which may lead to incorrect results and delay of results due to the need to reconfigure assay parameters.
3. When performing a backup on the ARCHITECT while the iARM is replenishing the wash buffer at the same time, the iARM loses communication with the System Control Center. The loss of communication may cause the wash buffer container to overflow. This has the potential to lead to physical and chemical hazards.

**Impact on
Patient Results**

Refer to **Appendix A** for details concerning any patient results impacted due to the issues identified in ARCHITECT Software version 9.41 and earlier.

**Necessary
Actions to be
Taken by
Customer**

Please follow the Necessary Actions required in **Appendix A** until software version 9.45 or software version 9.50 is installed.

Your Abbott representative will schedule a mandatory upgrade of your ARCHITECT Software version 9.45 or ARCHITECT Software version 9.50 depending on your system configuration.

If you have forwarded the product listed above to other laboratories, please inform them of this Product Correction and provide to them a copy of this letter.

Please retain this letter for your laboratory records.

**Contact
Information**

If you or any of the health care providers you serve have questions regarding this information, U.S. Customers please contact Customer Service at 1-877-4ABBOTT (available 24 hours a day, 7 days a week). Customers outside the U.S., please contact your local area Customer Service.

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online (<http://www.fda.gov/MedWatch/report.htm>), by mail (<http://www.fda.gov/MedWatch/getforms.htm>), by phone (1-800-332-1088), or by fax (1-800-FDA-0178).

If you have experienced any patient or user injury associated with this Field Action, please immediately report the event to your local area Customer Service.
