



# Recall

illumina®

Thursday, January 16, 2025

Quality Notification  
**Urgent Medical Device Recall**  
**Urgent Field Safety Notice**

Dear Customer,

Illumina is contacting you regarding a security vulnerability identified in the Baseboard Management Controller (BMC) in the DRAGEN™ Server for the NovaSeq™ 6000Dx instruments and Illumina DRAGEN Server for NextSeq™ 550Dx instruments. Affected servers are specified in Table 1 below. This notice outlines the issue summary, Illumina actions, and required customer actions. Please note, this security vulnerability is limited to these Illumina DRAGEN Servers. Other versions of DRAGEN Servers are not impacted.

**Table 1: Affected Products**

Product Affected	Catalog Number	Unique Device Identifier - Device Identifier Number
NovaSeq 6000Dx Instrument including DRAGEN Server	20068232	00816270020637
Illumina DRAGEN Server for NextSeq 550Dx	20086130	N/A

### Issue Summary

The Illumina Product Security Team recently identified an uncontrolled product security risk associated with the Baseboard Management Controller (BMC) used in the DRAGEN Servers listed above. The security vulnerability is due to configurations in remote network management and monitoring.

Illumina has determined that if an unauthorized actor were to exploit this vulnerability the actor could access the BMC remote console without providing a password (authentication bypass). Exploitation of this vulnerability could lead to unauthorized system access, manipulation of hardware settings, remote code execution, remote control, data breaches and denial of service.

At this time, Illumina has not received any reports and has no evidence indicating that this vulnerability has been exploited.

**Technical Support:**  
[techsupport@illumina.com](mailto:techsupport@illumina.com)

**Customer Care:**  
[customercare@illumina.com](mailto:customercare@illumina.com)

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For In Vitro Diagnostic Use. Not available in all countries or regions.



## Illumina Actions

Illumina is providing customers with instructions below on how to secure their server(s) and protect their system(s) against exploitation of this vulnerability, which when applied will remediate the vulnerability.

Failure to follow these instructions could leave your system exposed to the risks described above.

The pertinent local and international regulatory bodies, including the Competent Authorities, are being notified of this issue.

## Required Customer Actions

For all affected DRAGEN Servers listed in Table 1 above, please take the following actions to remediate this critical security vulnerability:

- Download and follow the Customer Instructions from [this](#) webpage. The full hyperlink is below:  
<https://support.illumina.com/support-content/bmc.html>

**Complete and return the Verification Form after carrying out all the steps in the instructions provided on your specific server(s) identified as affected in Table 1.**

**NOTE:** If you suspect your server may have been compromised by an unauthorized actor, please immediately unplug the network cable and contact Illumina Technical Support at [techsupport@illumina.com](mailto:techsupport@illumina.com).

If you experience an adverse event due to this security vulnerability with the use of any of the affected servers, please report it to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax. You can complete and submit the report online at [www.fda.gov/medwatch/report.htm](http://www.fda.gov/medwatch/report.htm). In regions outside the USA, please contact your local regulatory authority.

Illumina takes security issues very seriously. We are committed to supporting you in addressing this vulnerability. For any other questions or assistance, please contact [techsupport@illumina.com](mailto:techsupport@illumina.com). You may also be contacted by an external vendor on behalf of Illumina to ensure that you have the support you need.

Sincerely,

*Gary Workman*  
Electronically signed by:  
Gary Workman  
Reason: Approver  
Date: Jan 3, 2025 13:04  
EST

**Gary Workman**  
Vice President of Global Quality

*Karen Gutekunst*  
Electronically signed by:  
Karen Gutekunst  
Reason: Approver  
Date: Jan 2, 2025 11:39  
PST

**Karen Gutekunst**  
Vice President of Regulatory Affairs

**Technical Support:**  
[techsupport@illumina.com](mailto:techsupport@illumina.com)

**Customer Care:**  
[customercare@illumina.com](mailto:customercare@illumina.com)

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### Why You're Receiving This Notification

You are receiving this notification because our records indicate that you are the appropriate contact for your organization for product changes, product obsolescence, and quality issues.

Please be aware that these notifications contain essential information about our products and are not marketing communications. As such, you may receive these notifications despite having opted-out of receiving marketing communications from Illumina. If you are not the appropriate individual in your organization to receive these notifications, you may unsubscribe from these notifications by [submitting this form](#). For more information, please see our [Privacy Policy](#).

**Technical Support:**

[techsupport@illumina.com](mailto:techsupport@illumina.com)

**Customer Care:**

[customercare@illumina.com](mailto:customercare@illumina.com)

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### Verification Form

Dear Customer,

Illumina sent you an Urgent Medical Device Recall Notice FSN2025-1682 regarding an issue affecting the DRAGEN Servers for the NovaSeq 6000Dx and NextSeq 550Dx instruments.

Please complete the form below to confirm that you have received the notice and completed the Required Customer Actions outlined in the notification. Once completed, please email the form to [techsupport@illumina.com](mailto:techsupport@illumina.com).

Alternatively, you may e-mail Illumina Technical Support to provide the information requested below.

Verification Form	
<b>Company Name</b>	
<b>Information of Person Completing Form</b>	
Name:	
Title:	
Date (DD-MMM-YYYY):	
<b>Server Name</b>	<b>Serial Number(s)</b>
DRAGEN Server for NovaSeq 6000Dx	
Illumina DRAGEN Server for NextSeq 550Dx	
<b>Customer Responses</b>	
I confirm receipt of FSN2025-1682 and that I read and understood its content.	<input type="checkbox"/> Yes <input type="checkbox"/> No
The information has been brought to the attention of all relevant users.	<input type="checkbox"/> Yes <input type="checkbox"/> No
I confirm that the remediation steps listed in the Required Customer Actions section have been implemented on each and every affected server identified in this verification form.	<input type="checkbox"/> Yes <input type="checkbox"/> No

**Technical Support:**  
[techsupport@illumina.com](mailto:techsupport@illumina.com)

**Customer Care:**  
[customercare@illumina.com](mailto:customercare@illumina.com)

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Distributor/Importer Responses	<input type="checkbox"/> Not applicable
I have identified customers that received or may have received the affected servers.	<input type="checkbox"/> Yes <input type="checkbox"/> No
I have informed the identified customers of this recall.	<input type="checkbox"/> Yes <input type="checkbox"/> No Date (DD-MMM-YYYY):

**Technical Support:**  
[techsupport@illumina.com](mailto:techsupport@illumina.com)

**Customer Care:**  
[customercare@illumina.com](mailto:customercare@illumina.com)

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