

**RANDOX**  
Urgent Field Safety Notice

**Liquid Assayed Specific Protein Controls**

**Date:** 03 May 18

**Complaint Reference:** REC326

**Action Type:** Device Modification

**Detail on Affected Devices:**

Our records indicate that your facility may have received the following product.

Assay	Catalogue Number	GTIN
Liquid Assayed Specific Protein Controls	PS2682	05055273204896
	PS2683	05055273204902
	PS2684	05055273204919

**Reason for Recall:**

Randox have confirmed that Free Kappa light chains in our Liquid Assayed Specific Protein Controls increase over shelf life of the product.

**Risk to Health:**

The quality control results which are not within range can lead to a delay in reporting Free Kappa Light Chains results.

Free kappa light chains are used as an aid in diagnosis of monoclonal gammopathy of undetermined significance and for Multiple Myeloma through review of the ratio with free lamda chains. A short delay in reporting this result would not prevent diagnosis of these progressive diseases and therefore is unlikely to pose an immediate risk to health.

**Action to be taken:**

- Discuss the contents of this notice with your Medical Director.
- Place a copy of the important notice and updated IFU into any remaining stock.
- Complete and return the vigilance response section of this form to [technical.services@randox.com](mailto:technical.services@randox.com) within five working days.)

# **RANDOX**

## **Urgent Field Safety Notice**

### **Liquid Assayed Specific Protein Controls**


**Transmission of Field Safety Notice:** Send a copy of the FSN to all affected customers and to those who need to be aware within your organisation.

**Contact Reference:**

Radox Technical Services  
Radox Laboratories Ltd,  
55 Diamond Road,  
Crumlin,  
United Kingdom,  
BT29 4QY  
Email: [technical.services@radox.com](mailto:technical.services@radox.com)  
Tel: +44 (0) 28 9445 1070  
Fax: +44 (0) 28 9445 2912

Please accept our apologies for any inconvenience caused. Thank you for your patience and understanding. If you have any questions or concerns please contact Radox Technical Services.

**The undersigned confirms that this notice has been notified to the appropriate Regulatory Agency**

  
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**Vigilance Response Form** (Response Plan must be completed by the importer of the device)

**Importer Details**

Company Name	
Address	

**Total Quantity**

Received	
Distributed	

**Area of Distribution**

(To be completed by Distributors and Randox Offices)

Consignee	Country	Quantity Received	Analyser Serial Number	Replacements Required

I have read and understood the Urgent Field Safety Notice. The actions to be taken are completed.

Completed By				Date	
Contact	Tel		Email		

