

## **GRMS Recall & Incident Notification Form**

As part of the contractual relationship with certificated sites, the site shall notify the certification body of:

- any impending prosecution or enforcement with respect to product safety or legality
- all product recalls
- adverse media or regulatory authority interest
- evidence of a significant public safety issue (e.g. food poisoning outbreak or customer injury)
- evidence of significant failings at the certificated site (e.g. fraud, corruption or significant malpractice)
- adverse public statements by a regulatory authority, NGO or major retailer
- significant public safety concerns bringing scheme owner or BVC into disrepute

This contractual requirement is also reflected in the Standards (e.g. **notification to the certification body within 3** days).

The aim of this notification is to allow the certification body to assess whether the incident is indicative of a failure of the site's systems. The Certification Body must take the necessary steps to fully understand the implications of the situation and take appropriate actions. This may include requests for additional information, a further visit to the site, further full or partial re-audits, suspension or withdrawal of the certificate.

**Initial notification to scheme owner must be made within 24 hours of the site notifying the certification body.** A further update can be made, where necessary, to confirm the root cause and extend as well as the immediate corrections and subsequent corrective actions within a further 3 weeks.

## PLEASE FILL IN AND SENT THIS FORM TO: recalls@bureauveritas.com

SECTION I. To be completed by affected certificated site			
	l	illicated Site	
Name, phone and e-			
mail of responsible			
person at site notifying			
BVCDK of recall /			
incident			
Date of notification			
Site Code (Not mandatory			
to be filled in by site)			
Company/Site Name			
As it appears on the			
certificate			
Country			
Where the site is based		T	T
Certificate information	Certificate no.	Accreditation:	Validity:
Reason for notification			
Select one			
Category of Product			
Recall			
Select one			
Outline of			
Recall/Incident			
Briefly explain the reason			
for the incident or recall.			
Include if required by			
authority.			
Authority informed and			
when Did the recall or incident			
generate significant media			
coverage			
Has product reached			
consumer (Yes/No)			
Has there been any			
hospitalization or			
deaths? (Explain)			
ucatiis: (Explain)			



## **GRMS Recall & Incident Notification Form**

Product(s) effected	
Detail product name, type	
of product, batch codes	
effected if known otherwise	
update within 3 weeks	
Date of Recall or	
Incident	
What date the incident or	
recall start	
Extend and Correction	
(action taken by Site)	
Evaluated extend and	
action(s) taken by the site to	
rectify the incident/product	
Site or Supplier Issue	
Select one	
Product handling	
(returns, destruction)	
And	
% of product not	
accounted for	
Root Cause Analysis	
(conducted by Site) –	
If root cause cannot be	
confirmed immediately it	
must be reviewed and	
provided to Certification	
Body within 3 weeks of the	
date of recall.	
<b>Corrective Action Plan</b>	
(conducted by Site)	
(conducted by Site) If corrective action plan	
(conducted by Site) If corrective action plan cannot be confirmed	
If corrective action plan	
If corrective action plan cannot be confirmed immediately it must be reviewed and provided to	
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## **GRMS Recall & Incident Notification Form**

Date of Certification	
Status Change (If	
applicable)	
Date of suspension or	
withdrawal	
Any other information	

Notes: