

General Product 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				
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				
Certificate information	Certificate no.	Accreditation:	Validity:	
Reason for notification Select one		1		
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)				



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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.	
Corrective Action Plan	
(conducted by Site)	
IC	
If corrective action plan	
If corrective action plan cannot be confirmed	
If corrective action plan cannot be confirmed immediately it must be	
If corrective action plan cannot be confirmed immediately it must be reviewed and provided to	
If corrective action plan cannot be confirmed immediately it must be reviewed and provided to Certification Body within 3	
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Date of Certification Status Change (If
applicable)
Date of suspension or withdrawal
Any other information

Notes: