

BRCGS Product Safety Incidents Notification Form

Product safety incidents defined as an event that has occurred that may result in the production or supply of unsafe, illegal, or non-conforming products are required to be reported to protect the integrity and reputation of BRCGS, partner brands and specifiers, certification bodies, auditors, and sites and ensure appropriate action is taken by the site, including root cause analysis and preventive action.

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

- legal proceedings with respect to product safety or legality, or that which significantly affects the operation of the site
- enforcement by authorities related to product safety or legality (e.g. an enforcement notice)
- product recalls, food safety-related product withdrawals, any significant public food safety incidents, or any significant regulatory food safety non-conformities
- significant damage to the site (e.g. natural disaster such as flood or damage by fire)
- change of ownership (when the title is transferred from one individual or entity to another and results in a change of control of the organisation).
- any significant change to the operation or scope
- significant staff changes or prolonged shutdowns (e.g. considerable staff losses or the loss of key product safety roles).

This contractual requirement is also reflected in the Standards (e.g. BRCGS Food Safety Issue 9 clause 3.11.4 requires **notification to the certification body within 3 days** and general protocol requirements 6.1 Communication with certification bodies).

Sites certificated to the BRCGS Gluten-Free Certification Program must notify BRCGS at compliance@brcgs.com, where applicable the AOECS member and the Certification Body within 1 working day of the recall date.

The site shall then provide sufficient information to enable the certification body to assess any effects of the incident on the ongoing validity of the current certificate within **21 calendar days**. As a minimum, this shall include corrective action, root cause analysis and a preventive action plan.

The certification body in turn shall take appropriate steps to assess the situation and any implications for the certification, and shall take appropriate action. This may include requesting further details of the corrective action, root cause analysis and preventive action plan implemented by the site, undertake a site visit to verify the control of processes and confirm continued certification, suspension or withdrawal of the BRCGS certificate.

In the event of an incident, the effectiveness of corrective and preventive actions taken by the site will also be reviewed at the next scheduled BRCGS audit to confirm their implementation and continued effectiveness.

Changes to the certification status of a site shall be recorded in the BRCGS Directory.



BRCGS Product Safety Incidents Notification Form

SECTION A – To be completed by the BRCGS certificated site

Company / Site Name	
As listed in the BRCGS Directory	
BRCGS Site Code	
BRCGS Standard	
Reason for Notification	
Product recall : Any measure aimed at achieving the return of an unsafe or illegal product from a customer and consumer.	
Food safety-related withdrawal : Any measure aimed at achieving the return of an unsafe or illegal product from a customer.	
Regulatory notice: non-conformity raised by the regulator's official.	
Incident Category	
Where the risk is identified as an undeclared allergen, whether the recall is due to incorrect labelling, incorrect packaging or contamination of the product by an allergen these should all be listed under 'allergen'.	
Outline of Incident	
Briefly explain the reason for the incident.	
Products Recalled	
Product name and description, relevant to all incidents, not only recalls.	
Please use simple descriptions e.g. 'ready meal', chocolate', not the brand names and always provide a product description when the product name is not provided in English.	
Date of Recall	Select a date
Date when the incident was started at the site	Colour a date
Date when the incident was started at the site	
Correction (action taken by the site)	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility.	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility.	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility. Site or Supplier	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility. Site or Supplier Specify the identified source of the incident	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility. Site or Supplier Specify the identified source of the incident Root Cause Analysis (within 21 days of incident) Identify the underlying cause of the recall/incident except for some traded goods or Storage and Distribution sites where the following	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility. Site or Supplier Specify the identified source of the incident Root Cause Analysis (within 21 days of incident) Identify the underlying cause of the recall/incident except for some traded goods or Storage and Distribution sites where the following actions would be required: For Storage and Distribution sites, where supplier approval is not a part of the scope of the certification, and the cause of the incident does not involve any action by the site, the root cause analysis may not be within	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility. Site or Supplier Specify the identified source of the incident Root Cause Analysis (within 21 days of incident) Identify the underlying cause of the recall/incident except for some traded goods or Storage and Distribution sites where the following actions would be required: For Storage and Distribution sites, where supplier approval is not a part of the scope of the certification, and the cause of the incident does not involve any action by the site, the root cause analysis may not be within the scope of the site operation.	
Correction (action taken by the site) Outline the steps taken immediately by the site covering their scope of responsibility. Site or Supplier Specify the identified source of the incident Root Cause Analysis (within 21 days of incident) Identify the underlying cause of the recall/incident except for some traded goods or Storage and Distribution sites where the following actions would be required: For Storage and Distribution sites, where supplier approval is not a part of the scope of the certification, and the cause of the incident does not involve any action by the site, the root cause analysis may not be within the scope of the site operation. Or In some instances, supplier actions would be required. Where Traded Goods or Storage & Distribution site failure results in an incident, a full investigation is required. Storage and Distribution sites shall undertake	



BRCGS Product Safety Incidents Notification Form

Details of who completed this form

Contact Name	
Contact Position	
Contact Phone Number	
Contact E-mail Address	
Date	
Signed	

SECTION B – To be completed by the Certification Body Local Office

Date of Local Office Submission to ICC UK (24 Hours)	
	Select a date
Date completed form submitted to brcgsrecalls@bureauveritas.com	
Local Office Contact & Location	
BV contact and office managing the communication with the site	
Additional Information	
Late site /local office notifications and action taken. Should further details	
be pending (e.g. RCA, Microbiological test results) when these become	
available, an update should be provided.	

SECTION C - To be completed by ICC UK

BRCGS Submission ID	
BRCGS Submission By	
BRCGS Initial Submission Date	
	Select a date
(within 2 working days of site notification)	
Certification Status	
CB to confirm if the site certification was affected. Use "Certificated" for the sites whose certification status was not affected.	
Date of Certification Status Change	
	Select a date
If applicable	
Final Submission Date	
Date when all information related to the incident (including RCA and PAP) has been completed and submitted to BRCGS for approval – the information should be provided within 23 calendar days from the date of recall.	Select a date
Additional Information	
Justification for changed in certification status.	
Recall Status	