CMAP Supplier Campaigning

-

Supplier's Quick Reference Guide

Rev 3

September 18th, 2026

EVERY CONNECTION COUNTS

Overview of CMAP Supplier Campaigning Quick Guide

TE Connectivity is engaging its supply chain to collect information regarding legislation on restricted and reportable substances in products. TE has selected the **Compliance Map (CMAP)** platform to support this initiative. This reference guide provides helpful resources and instructions to assist you in submitting your responses.

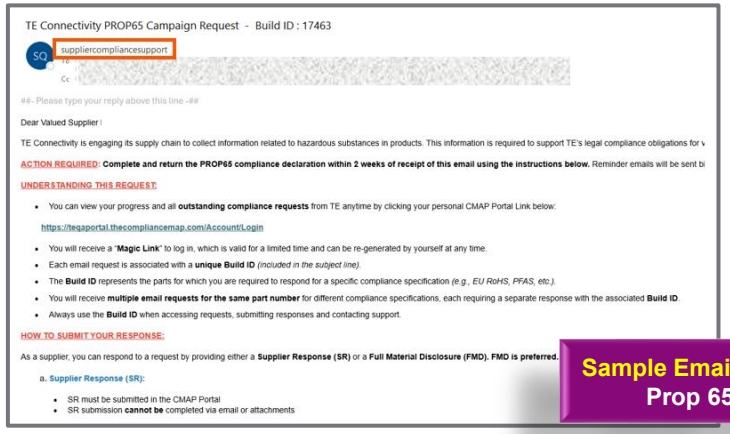
This Quick Guide covers how suppliers:

- **Receive campaign notifications**
- **Access CMAP Portal**
- **Complete and submit required surveys and declarations for compliance specs like EU ROHS, EU REACH, PFAS etc.**
- **Complete TE Medical Device Survey (TE custom survey)**
- **Conflict Mineral Reporting Template (CMRT)**
- **Extended Mineral Reporting Template (EMRT)**

Legends & Descriptions

Abbreviation	Definition
CMAP	The Compliance Map
SR	**Supplier Response** (a high-level compliance declaration submitted by a supplier in the CMAP Portal to confirm whether a part contains or does not contain regulated substances under a specific compliance requirement to confirm the compliance status of a part)
FMD	**Full Material Disclosure** (a detailed declaration submitted by a supplier that provides the complete material composition of a part, including all materials and substances present, along with their respective quantities).
RoHS	Restriction of Hazardous Substances
REACH	Registration, Evaluation, Authorization and Restriction of Chemicals
CMRT	Conflict Minerals Reporting Template
EMRT	Extended Minerals Reporting Template
PFAS	Per- and Polyfluoroalkyl Substances
Class C	The form type to be selected in the CMAP Portal for submitting Supplier Response (SR)
TE MD	TE Medical Device Survey (TE'S custom campaign)

The Compliance Specs & Campaign Requests


TE Connectivity PROP65 Campaign Request - Build ID : 17463

suppliercompliancesupport

Dear Valued Supplier !

TE Connectivity is engaging its supply chain to collect information related to hazardous substances in products. This information is required to support TE's legal compliance obligations for v

ACTION REQUIRED: **Complete and return the PROP65 compliance declaration within 2 weeks of receipt of this email using the instructions below.** Reminder emails will be sent bi

UNDERSTANDING THIS REQUEST:

- You can view your progress and all **outstanding compliance requests** from TE anytime by clicking your personal CMAP Portal Link below:
  https://teqaportal.thecompliancemap.com/Account/Login
- You will receive a "**Magic Link**" to log in, which is valid for a limited time and can be re-generated by yourself at any time.
- Each email request is associated with a **unique Build ID** *(included in the subject line)*.
- The **Build ID** represents the parts for which you are required to respond for a specific compliance specification *(e.g., EU RoHS, PFAS, etc.)*.
- You will receive **multiple email requests for the same part number** for different compliance specifications, each requiring a separate response with the associated **Build ID**.
- Always use the **Build ID** when accessing requests, submitting responses and contacting support.

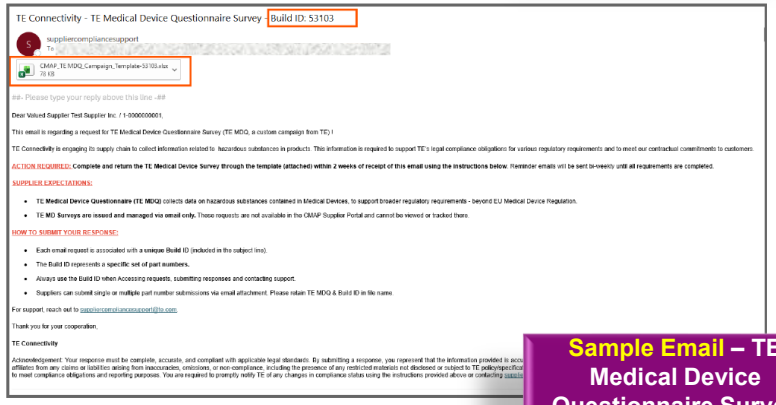
HOW TO SUBMIT YOUR RESPONSE:

As a supplier, you can respond to a request by providing either a **Supplier Response (SR)** or a **Full Material Disclosure (FMD). FMD is preferred.**

a. **Supplier Response (SR):**

- SR must be submitted in the CMAP Portal
- SR submission **cannot be** completed via email or attachments

Sample Email – CA Prop 65


TE Connectivity - TE Medical Device Questionnaire Survey Build ID: 53103

suppliercompliancesupport

CMAP_TE.MDQ_Campaign_Template-53103.xlsx 20 KB

Dear Valued Supplier Test Supplier Inc. / 1-0000000001,

This email is regarding a request for TE Medical Device Questionnaire Survey (TE MDQ, a custom campaign from TE) !

TE Connectivity is engaging its supply chain to collect information related to hazardous substances in products. This information is required to support TE's legal compliance obligations for various regulatory requirements and to meet our contractual commitments to customers.

ACTION REQUIRED: **Complete and return the TE Medical Device Survey through the template (attached) within 2 weeks of receipt of this email using the instructions below.** Reminder emails will be sent bi-weekly until all requirements are completed.

SUPPLIER EXPECTATIONS:

- TE **Medical Device Questionnaire (TE MDQ)** collects data on hazardous substances contained in Medical Devices, to support broader regulatory requirements - beyond EU Medical Device Regulation.
- **TE MD Surveys are issued and managed via email only.** These requests are not available in the CMAP Supplier Portal and cannot be viewed or tracked there.

HOW TO SUBMIT YOUR RESPONSE:

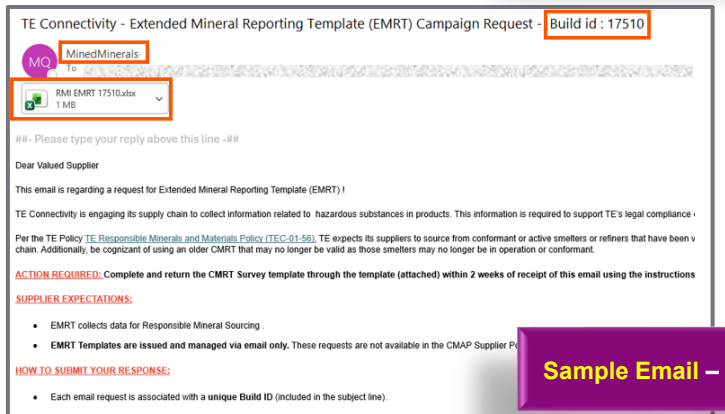
- Each email request is associated with a **unique Build ID** (included in the subject line).
- The Build ID represents a specific set of part numbers.
- Always use the Build ID when Accessing requests, submitting responses and contacting support.
- Suppliers can submit single or multiple part number submissions via email attachment. Please retain TE MDQ & Build ID in file name.

For support, reach out to suppliercompliancesupport@te.com

Thank you for your cooperation,

TE Connectivity

Sample Email – TE Medical Device Questionnaire Survey


TE Connectivity - Extended Mineral Reporting Template (EMRT) Campaign Request - Build id : 17510

MinedMinerals

RMI EMRT 17510.xlsx 1 MB

Dear Valued Supplier

This email is regarding a request for Extended Mineral Reporting Template (EMRT) !

TE Connectivity is engaging its supply chain to collect information related to hazardous substances in products. This information is required to support TE's legal compliance

Per the TE Policy TE Responsible Minerals and Materials Policy (TEC-01-56), TE expects its suppliers to source from conformant or active smelters or refiners that have been v chain. Additionally, be cognizant of using an older CMRT that may no longer be valid as those smelters may no longer be in operation or conformant.

ACTION REQUIRED: **Complete and return the CMRT Survey template through the template (attached) within 2 weeks of receipt of this email using the instructions**

SUPPLIER EXPECTATIONS:

- EMRT collects data for Responsible Mineral Sourcing .
- **EMRT Templates are issued and managed via email only.** These requests are not available in the CMAP Supplier P

HOW TO SUBMIT YOUR RESPONSE:

- Each email request is associated with a **unique Build ID** (included in the subject line).

Sample Email – EMRT

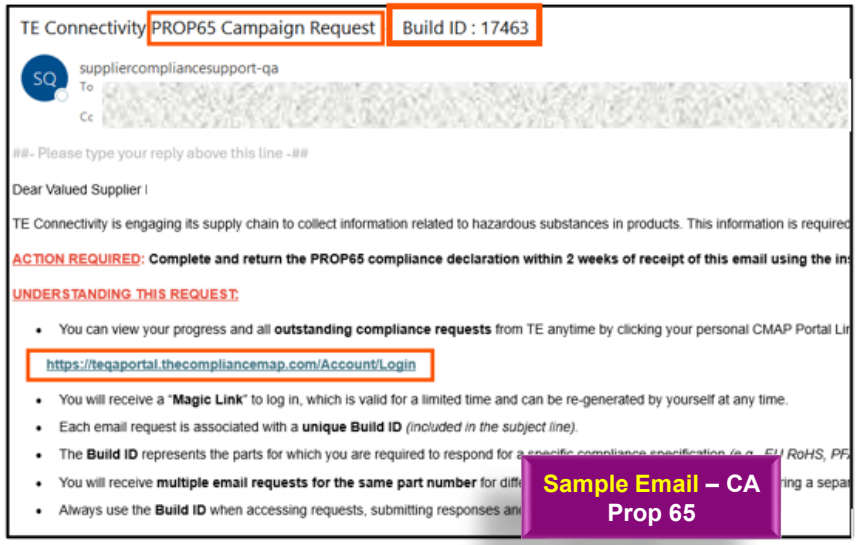
Suppliers will receive campaign requests for the below compliance specs from (suppliercompliancesupport@te.com):

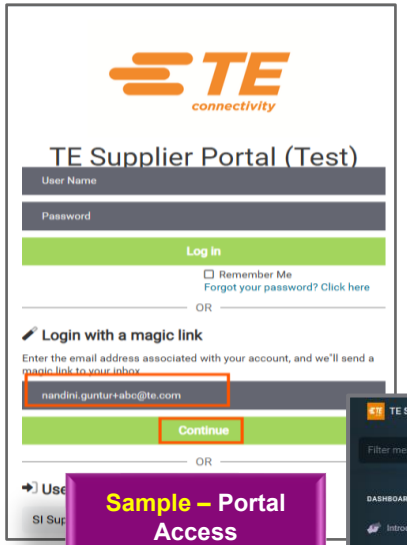
- **EU RoHS**
- **EU REACH SVHC**
- **EU REACH Annex XVII (Article 67 / Restriction List)**
- **CA Proposition 65**
- **PFAS**
- **TE Low Halogen**
- **TE Banned & Targeted Substance**
- **TE Medical Device Questionnaire Survey** *(with TE Custom Medical Device Template attached in campaign email)*

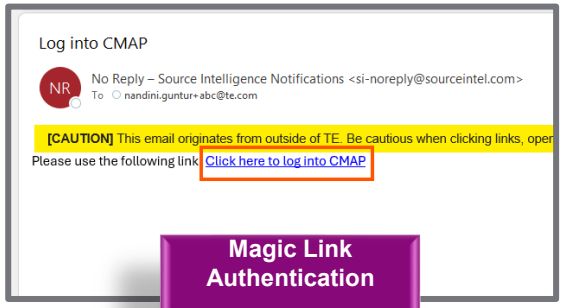
Campaign requests for the below will be received from minedminerals@te.com *(with the respective templates attached in campaign email including Build ID in file name)*

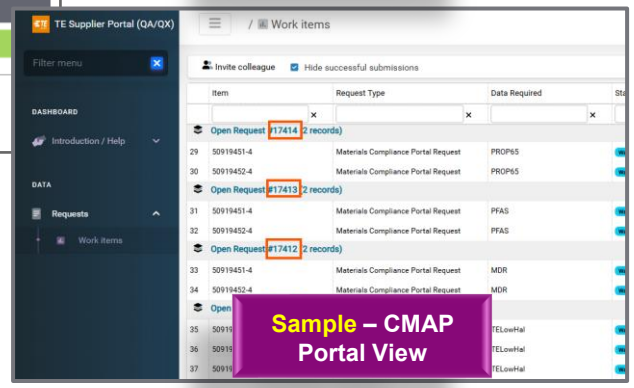
- **Conflict Mineral Reporting Template (CMRT)**
- **Extended Mineral Reporting Template (EMRT)**

Key Concepts: Portal Access & Build ID Management and









Email Notification Access

Access starts with a campaign notification email containing the portal link sent to the registered supplier email address(es).

Magic Link Authentication

Suppliers receive a secure, Magic Link via email to log in without passwords sent to the registered email for time-limited access authentication.

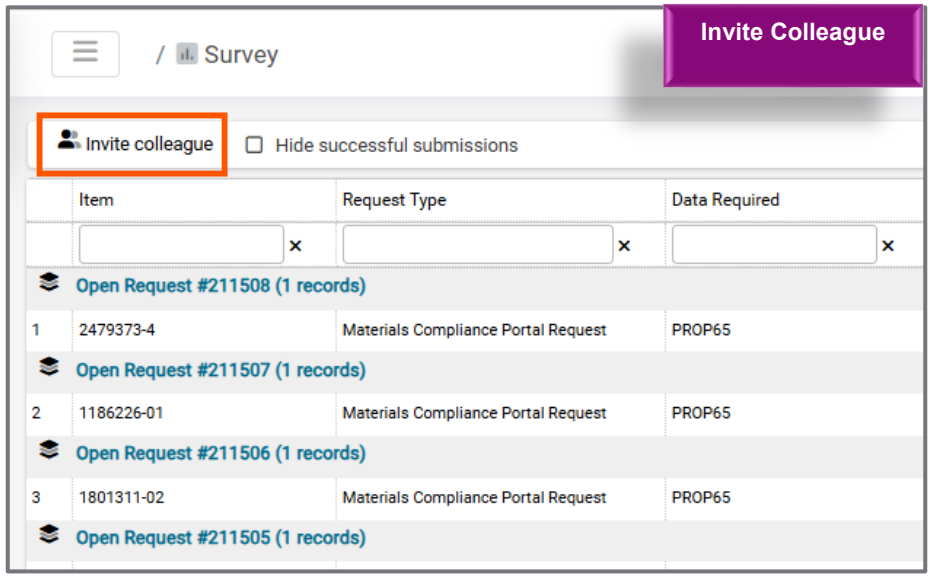
Unique Build ID

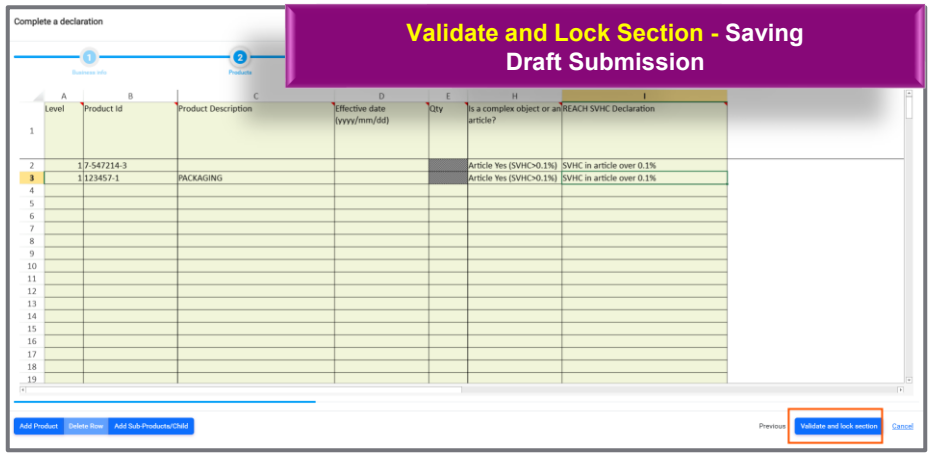
Build ID uniquely identifies each compliance campaign, linking part numbers to each compliance specification. This will be indicated in the subject of the email and on the CMAP Portal.

Build Id Generation - Campaign Consolidation

When TE issues a new campaign for the same Compliance Specification, any newly requested parts and outstanding (unanswered) parts from previous Build IDs will be consolidated into a new Build ID. Suppliers should always respond using the latest Build ID, as older Build IDs will be closed. Previously submitted parts will not be requested again, and only one active Build ID will be available at a time.

Key Concepts: Submission, Collaboration & Draft Saving





Submission Alignment

Submissions which are to be sent via email (TE Medical Device Survey/ CMRT/EMRT except IPC 1752A) must include Build ID number in the file name and be returned as attachment to the Build ID email.

Collaboration and Data Management

Suppliers can invite colleagues to contribute and manage multiple parts with copy and delete features via '**Invite Colleague**' button.

Saving A Draft Submission

Use the Validate and Lock Section button to save your responses as a draft on the portal. This allows you to save your progress and return later to complete or update your submission before final submission.

Types of Submissions

Supplier Responses (SR)

SRs are high-level declarations confirming the presence or absence of reportable substances under a specific regulation.

→ Use Class C in CMAP Portal to complete **Supplier Responses** for parts

→ Mandatory (**Red**) fields must be completed

→ **DO NOT** enter data in greyed out fields as these are not applicable to fill

→ Selecting the correct compliance status, such as compliant, compliant by exemption and non-compliant, is essential for accuracy.

Full Material Declarations (FMD)

FMDs provide detailed disclosure of all materials including weights and percentages in a part

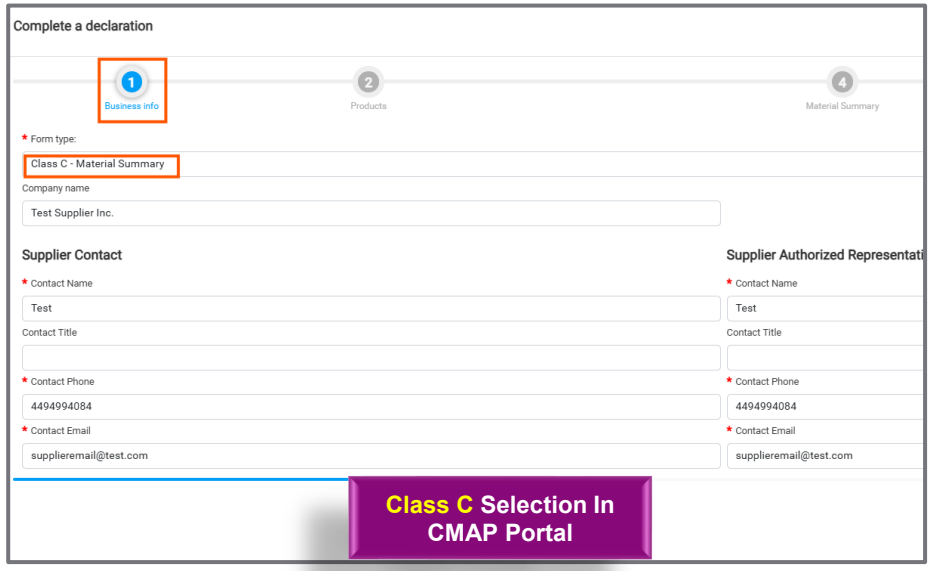
→ Suppliers can send the IPC 1752A XML files for each part number via email.

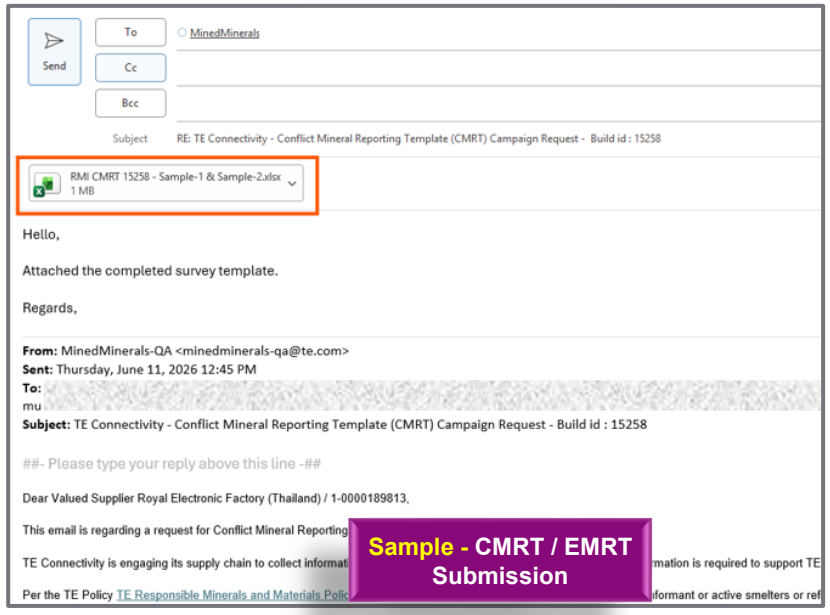
Responsible Minerals Reporting

CMRT and EMRT templates use standardized templates, completed offline and submitted via email.

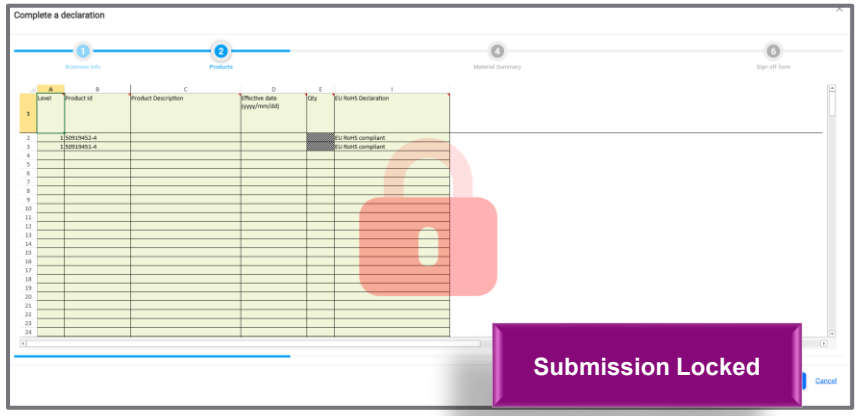
Custom TE Medical Device Survey Templates

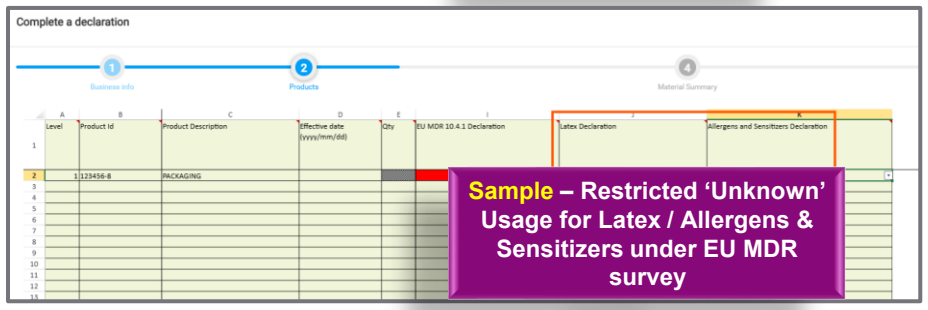
TE Medical Device Survey use custom templates sent by email instead of the CMAP Portal.

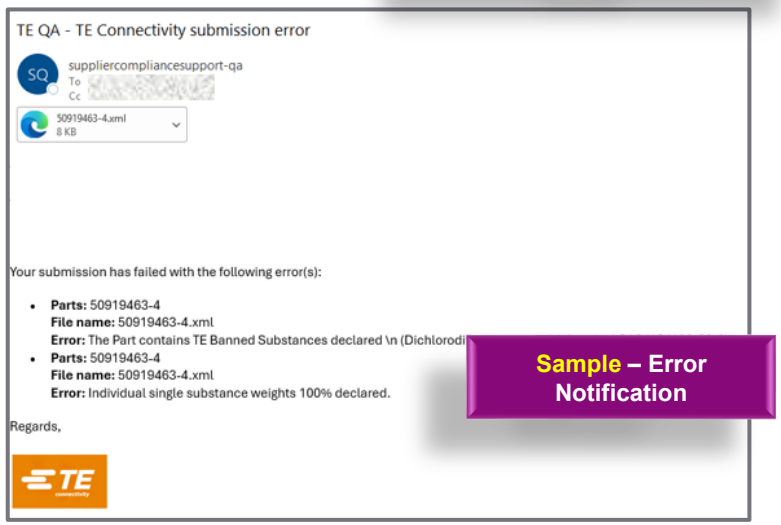




Important Rules and Controls







Submission Locking

Once an SR on a Build ID is submitted, the survey form will be locked and cannot be edited. This is to preserve data integrity.

Methods To Override Previous Submissions For Suppliers

If changes are required after submission, the following alternative options are available:

1.Submit a Full Material Declaration (FMD) in IPC 1752A XML format as attachment via email in response to the campaign notification email as guided in **Section-IV** of the supplier manual.

2. If supplier prefers to submit or override the older submission via a new Supplier Response (SR), send request to TE at suppliercompliancesupport@te.com to create a new campaign providing your old Build ID.

Once the new campaign is created, you can submit a new SR for the same compliance specification.

The new Supplier Response (SR) submission shall override the older submission in TE.

Restricted 'Unknown' Usage

Use of '**Unknown**' responses to any compliance specification is limited and only permitted for questionnaire on - **Latex or Allergens and Sensitizers Declaration under EU Medical Device Regulation**.

Notifications on Errors & Warnings

Suppliers may receive automated email notifications for errors or warnings. For example, if a submission has weight mismatches, they will be notified and required to correct and resubmit.

Key Takeaways

- **Correct Submission Channels**

  Always respond using specified channels like portal for **SR** and email for **FMD, CMRT, EMRT and TE MD** to ensure accurate communication.

- **Utilizing Portal Features**

  Use portal tools like filtering, colleague invitation, and status tracking to manage workload efficiently and timely.

- **Proactive Support Engagement**

  Ensure campaign emails are received and contact compliance support at suppliercompliancesupport@te.com proactively to resolve issues quickly.