FAILURE REPORTING, ANALYSIS AND CORRECTIVE ACTION SYSTEM

FRACAS

A PARTNERING FOR TOTAL QUALITY DOCUMENT

SEMA TECH
FAILURE REPORTING, ANALYSIS, AND CORRECTIVE ACTION SYSTEM
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ACKNOWLEDGMENTS

In putting together this document, there were many obstacles that kept getting in the way of completing it. Without the support of my family, Sonia, Jason, and Christopher, this would never have been accomplished.

Mario Villacourt
External Total Quality and Reliability
SEMATECH

I would like to thank my wife, Monique I. Govil, for her support and encouragement in completing this work. Also, I want to thank SVGL for supporting my participation with this effort.

Pradeep Govil
Reliability Engineering
Silicon Valley Group Lithography Systems
Failure Reporting, Analysis, and Corrective Action System (FRA-CAS) is a closed-loop feedback path in which the user and the supplier work together to collect, record, and analyze failures of both hardware and software data sets. The user captures predetermined types of data about all problems with a particular tool or software and submits the data to that supplier. A Failure Review Board (FRB) at the supplier site analyzes the failures, considering such factors as time, money, and engineering personnel. The resulting analysis identifies corrective actions that should be implemented and verified to prevent failures from recurring. A simplified version of this process is depicted in Figure 1.
Since factors vary among installation sites, equipment users must work closely with each of their suppliers to ensure that proper data is being collected, that the data is being provided to the correct supplier, and that the resulting solutions are feasible.

Unlike other reliability activities, FRACAS promotes reliability improvement throughout the life cycle of the equipment. The method can be used during in-house (laboratory) tests, field (alpha or beta site) tests, and production/operations to determine where problems are concentrated within the design of the equipment.

According to the military,

> Corrective action options and flexibility are greatest during design evolution, when even major design changes can be considered to eliminate or significantly reduce susceptibility to known failure causes. These options and flexibility become more limited and expensive to implement as a design becomes firm. The earlier the failure cause is identified and positive corrective action implemented, the sooner both the producer and user realize the benefits of reduced failure occurrences in the factory and in the field. Early implementation of corrective action also has the advantage of providing visibility of the adequacy of the corrective action in the event more effort is required.

In addition to its military use, FRACAS has been applied extensively in the aerospace, automotive, and telecommunications industries. Elements of FRACAS can be found throughout all industries, but this limited use usually means that the only data received by the manufacturer are complaints logged with the field service or customer service organizations.

FRACAS can provide control and management visibility for improving the reliability and maintainability of semiconductor manufacturing equipment hardware and software. Timely disciplined use of failure and maintenance data can help generate and implement effective corrective actions to prevent failures from recurring and to simplify or reduce maintenance tasks.

FRACAS objectives include the following:

- Providing engineering data for corrective action
- Assessing historical reliability performance, such as mean time between failures (MTBF), mean time to repair (MTTR), availability, preventive maintenance, etc.
- Developing patterns for deficiencies
- Providing data for statistical analysis

Also, FRACAS information can help measure contractual performance to better determine warranty information.

FRACAS provides a complete reporting, analysis, and corrective action process that fulfills ISO 9000 requirements. More and more companies are requiring their suppliers to meet ISO 9000, for example, the SEMATECH Standardized Supplier Quality Assessment (SSQA) and the Motorola Quality Systems Review (QSR).

**FAILURE REPORTING**

All events (failures) that occur during inspections and tests should be reported through an established procedure that includes collecting and recording corrective maintenance information and times. The data included in these reports should be verified and then the data should be submitted on simple, easy-to-use forms that are tailored to the respective equipment or software.
Collecting the Data

Many problems go unnoticed because insufficient information was provided. The FRB must know if, for example, someone was able to duplicate the problem being reported. There are three common causes for missing essential data:

- Inspection or testing began before a procedure was in place to report problems.
- The reporting form was difficult to use.
- The person who filled out the form had not been trained.

Operators and maintenance personnel are usually the first to identify problems and, therefore, they should be trained to properly capture all of the information needed for an event report. The contents of this report are more fully described in the next section.

However data is collected, one constant should be emphasized: consolidate all the data into a central data logging system. When field failures occur, two types of data collection are usually present: (1) the user’s system that includes equipment utilization and process-related information, and (2) field service reports (FSRs) that include parts replacement information. The supplier has great difficulty addressing problems when FSRs are the only source of data. It is helpful if the user provides the supplier with detailed logs for problem verification. The supplier should correlate both types to help determine priorities for problems to be resolved.

The supplier should be included in the formation of the user’s data collection system while the product is being installed. Ideally, the equipment should log all information automatically and then download the data to the supplier’s data collection system. This would eliminate the need for paper forms and also the confusion caused by duplicate data sets.
Reporting Equipment Failures

Collecting and sharing appropriate data through event reports are essential components of an effective FRACAS process, both for the supplier and for the user. There are common elements in every report (when the event occurred, what item failed, etc.) that the user and the supplier both use to analyze failures. Other crucial information includes the duration of the failure, the time it took to repair, and the type of metric used (time or wafer cycles).

To simplify the effort of collecting data and to minimize any duplication of effort, use the following guidelines:

- Capture data at the time of the failure.
- Capture utilization data through the user’s factory tracking system (for example, WorkStream).
- Avoid duplicating data collection between the supplier’s and the user’s systems.
- Make failure and utilization data available to both the supplier and the user in a standardized format.
- Exchange failure and utilization data between the supplier and user frequently.
- Link event reports to the corresponding utilization data in the factory tracking system through an identifying event number that is captured on the equipment.
- Automate both the user’s and the supplier’s systems to maximize efficiency, minimize paper tracking, and avoid dual reporting.

In addition, make full use of reporting tools that already exist (to which little or no modification may be necessary), such as

- In-house test reports
- Field service reports
- Customer activity logs
- Customer equipment history

Any other pertinent information you may need may be determined by asking the design engineers what information they need for root cause analysis. Data log files (history kept by the software) should accompany the failure report to provide software engineers with information about events that preceded the malfunction.

As a result of the data collected, minimum reliability performance metrics (such as MTBF, MTTR, and uptime percentage) could be determined for each system.

Every piece of data helps. In addition to these common elements, there are specific types of data needed only by the user or only by the supplier.

The supplier should acquire detailed failure information from the event report that should become an integral part of the equipment log file. The details help determine the state of operation before and after the failure, which is necessary to address the root cause of equipment failures efficiently and effectively.

Table 1 shows the types of detail that the supplier needs to see in the event report (in-house or field). Figure 2 is an example of an event report form. The format of the form is important only to simplify the task of the data recorder. You may want to computerize your data entry forms to expedite the process and also minimize failure description errors.
<table>
<thead>
<tr>
<th><strong>Event Report Fields</strong></th>
<th><strong>Example</strong></th>
<th><strong>Description</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>Serial Number</td>
<td>001</td>
<td>System-unique identifier that indicates who the customer is and where the system is located</td>
</tr>
<tr>
<td>Date and Time of Event</td>
<td>March 12, 1994 at 4:00 PM</td>
<td></td>
</tr>
<tr>
<td>Duration of the Event (Downtime)</td>
<td>01:21</td>
<td>Usually in hours and minutes</td>
</tr>
<tr>
<td>Failed Item Description</td>
<td>Handler-Stock-Robot-Motor</td>
<td>Describes function and location of failed item from top to bottom; that is, field to lowest replaceable unit</td>
</tr>
<tr>
<td>Reliability/Fault Code of Known Problems</td>
<td>AH-STK-ROT-MOT-021</td>
<td>Known problem categorized by reliability code and failure mode number</td>
</tr>
<tr>
<td>Life Cycle Phase of the System</td>
<td>Field Operations</td>
<td>See <em>equipment life cycle</em> in the Glossary</td>
</tr>
<tr>
<td>Downtime Category (Maintenance Action)</td>
<td>unscheduled</td>
<td>Usually shown as either unscheduled or scheduled (preventive maintenance)</td>
</tr>
<tr>
<td>Part Number</td>
<td>244-PC</td>
<td>Part number of replaced item (if any)</td>
</tr>
<tr>
<td>Operator’s Name</td>
<td>M. Lolita</td>
<td>Person reporting problem</td>
</tr>
<tr>
<td>Service Engineer’s Name</td>
<td>S. Spade</td>
<td>Person performing repair action</td>
</tr>
<tr>
<td>Event Report Identification Number</td>
<td>FSR-002</td>
<td>Supplier’s report number</td>
</tr>
<tr>
<td>Event (Problem) Description</td>
<td>What happened?</td>
<td>Description of all conditions prior to failure and how observer thinks it failed</td>
</tr>
<tr>
<td>Repair Action Description</td>
<td>What did you do to repair item?</td>
<td>Description of any repair or maintenance attempts</td>
</tr>
</tbody>
</table>
**EVENT RECORD**

<table>
<thead>
<tr>
<th>Serial Number:</th>
<th>9003</th>
<th>Reliability Code:</th>
<th>DMD-XS-HRN-M3S-002</th>
</tr>
</thead>
<tbody>
<tr>
<td>Date:</td>
<td>09/01/93</td>
<td>Down Time Category:</td>
<td>Unscheduled</td>
</tr>
<tr>
<td>Time:</td>
<td>09:00:</td>
<td>Life Cycle Phase:</td>
<td>OPERATION</td>
</tr>
<tr>
<td>Duration:</td>
<td>04:30:</td>
<td>Relevant:</td>
<td>Y</td>
</tr>
<tr>
<td>System:</td>
<td>DMD</td>
<td>PM:</td>
<td>N</td>
</tr>
<tr>
<td>Subsystem:</td>
<td>XS</td>
<td>Part Number:</td>
<td>650-000029</td>
</tr>
<tr>
<td>Assembly:</td>
<td>HRN</td>
<td>Operator:</td>
<td>FRANK BELL</td>
</tr>
<tr>
<td>Subassembly:</td>
<td>M3S</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sub-subassembly:</td>
<td></td>
<td>FSE Name:</td>
<td></td>
</tr>
<tr>
<td>Problem:</td>
<td>PREVIOUS ATTEMPT TO RESOLVE TRANSPORT LOAD FAILURE DID NOT WORK. REQUIRES SWITCH REPLACEMENT.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Repair Action:</td>
<td>REPLACED SWITCH, CURED LENS 2, ALIGNED COLUMN, CHECKED JET ALIGN, MADE TEST MILLS AND DEPOS.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Figure 2* Example Event Report

The user analyzes this data to evaluate management of equipment and resources and to determine its effectiveness in meeting internal commitments. Internally, user data may be handled differently than described in SEMI E10-92, but as a minimum for sharing data with the supplier, the major states in Figure 3 should be followed.

*Figure 3* Equipment States Stack Chart
Reporting Software Problems

When reporting software problems, too much detail can be counterproductive. An error in software does not necessarily mean that there will be a problem at the system (equipment) level. To quote Sam Keene, 1992 IEEE Reliability Society president,

If I’m driving on a two-lane road (same direction) and switch lanes without using the signal indicator and continue to drive, I’ve committed an error. But if I do the same when another car is on the other side and I happen to crash into it, then I now have a problem.

Your existing software error reporting process should become an integral part of your overall FRACAS. (Rather than reiterating the software reliability data collection process, a practical approach can be found in A.M. Neufelder’s work.²)

FRACAS helps you focus on those errors that the customer may experience as problems. One focusing mechanism is to track the reason for a corrective action. This data can be collected by reviewing the repair action notes for each problem in the problem report. Engineering and management can categorize these reasons to determine the types of errors that occur most often and address them by improving procedures that most directly cause a particular type of error. The process of analyzing this data is continuous.

Among the reasons for corrective action are the following:

- Unclear requirements
- Misinterpreted requirements
- Changing requirements
- Documented design not coded
- Bad nesting
- Missing code

² Refer to Chapter 7, Software Reliability Data Collection, of Ensuring Software Reliability, by A. M. Neufelder (Marcel Dekker, Inc., New York, 1993).
- Excess code
- Previous maintenance
- Bad error handling
- Misuse of variables
- Conflicting parameters
- Software or hardware environment change
- Specification error
- Problem with third-party software
- Documented requirements not designed
- Software environment error (for example, an error in the compiler)

Reasons should also be ranked in terms of the criticality or severity of the error. This information helps management predict those errors that will cause downtime (as opposed to all errors, including those that may not cause downtime).

Severity rankings are:

1. Catastrophic
2. Severe
3. Moderate (has work-around)
4. Negligible
5. All others (for example, caused by a misunderstood new feature or unread documentation)

Also, tracking the modules or procedures modified for each corrective action helps schedule pre-release regression testing on these changes, which results in more efficient test procedures and more effective test results.
Logging Data

Whether software or equipment errors are being reported, once the supplier receives the reports, all data should be consolidated into one file. The supplier’s reliability/quality organization should oversee the data logging.

A FRACAS database is recommended for this part of the process. This is not a product-specific database but rather a database for tracking all products offered by a company. Figure 4 shows how the status of problems is kept updated in the database.

![PROBLEMS RECORD](attachment:image)

**Figure 4  Problems Record**
Using FRACAS Reports

The FRACAS database management system (DBMS) can display data in different ways. This section describes some of the report types you may find useful. A complete failure summary and other associated reports that this DBMS can provide are described in more detail beginning on page 27.

The graphical reports provide a quick snapshot of how the equipment or software is performing at any given point. Figure 5 shows the percentage rejection rate on a monthly basis as actual-versus-target rates. Figure 6 depicts the number of failure modes or problems on a weekly basis. Figure 7 points out the number of events being reported weekly by life cycle phase. Figure 8 provides a one-page status outline for tracking by the FRB. Whichever report you choose, it should be tailored to provide summaries and special reports for both management and engineering personnel.

Factory Test Tracking
Initial Test Reject Rate

![Graph](image)

**Figure 5** Example—Percentage Rejection Rate, Actual Versus Target

Failure Reporting, Analysis, and Corrective Action System
By Reliability Code
From 08/30/93 to 10/03/93

Figure 6  Example—Number of Events Weekly

Events by Life Cycle Phase by Week
From 09/27/93 to 10/17/93

Figure 7  Example—Number of Events Weekly by Life Cycle Phase
Figure 8 Example — Failure/Repair Report
Failure analysis is the process that determines the root cause of the failure. Each failure should be verified and then analyzed to the extent that you can identify the cause of the failure and any contributing factors. The methods used can range from a simple investigation of the circumstances surrounding the failure to a sophisticated laboratory analysis of all failed parts.

**Failure Analysis Process**

Failure analysis begins once an event report is written and sent to the FRB. It ends when you sufficiently understand the root cause so you can develop logically derived corrective actions.

It is important to clearly communicate the intent and structure of the failure analysis process to all appropriate organizations. These organizations should review and approve the process to confirm their cooperation. During and after the analysis, the problem owner, associated FRB member, and reliability coordinator must ensure that the database is maintained with the current applicable information.

The analysis process is:

1. Review, in detail, the field service reports.
2. Capture historical data from the database of any related or similar failures.
3. Assign owners for action items.
4. Do a root cause analysis (RCA).
5. Develop corrective actions.
6. Obtain the failed items for RCA (as needed).
7. Write a problem analysis report (PAR) and, if needed, a material disposition report (MDR).
Developing proper forms for tracking the failed parts and reporting the problem analysis results is essential. Figures 9 and 10 are examples of these types of forms.

<table>
<thead>
<tr>
<th>MATERIAL DISPOSITION REPORT</th>
</tr>
</thead>
<tbody>
<tr>
<td>S.NO.</td>
</tr>
<tr>
<td>J37511</td>
</tr>
</tbody>
</table>

- **RETURNED PART DESCRIPTION**
  - PART NUMBER
  - QTY RETURNED
  - PART DESCRIPTION
  - FSR NUMBER

- **REASON FOR RETURN**
  - DEFECTIVE
  - ENG CHANGE
  - OVERISSUE
  - OVERDRAWN
  - OTHER

- **USED ON**
  - EQUIPMENT S.NO.
  - ASSEMBLY NO.
  - SITE NUMBER

- **MATERIAL DISPOSITION INSTRUCTIONS**
  - DATE
  - RETURN TO STOCK
  - SEND TO FRB FOR FAILURE ANALYSIS
  - SEND FOR VENDOR FAILURE ANALYSIS
  - SCRAP
  - RETURN TO VENDOR
  - SEND FOR REPAIR

- **RELIABILITY INFORMATION**
  - TASK NUMBER
  - FRB MEMBER

<table>
<thead>
<tr>
<th>ORIGINATOR</th>
<th>DATE</th>
</tr>
</thead>
</table>

**DEFECT DESCRIPTION (FILLED BY THE ORIGINATOR)**

- **DISTRIBUTION**
  - RELIABILITY
  - MDC
  - FS
  - QUALITY

*Figure 9  MDR Forms*
PROBLEM ANALYSIS REPORT

PROBLEM NUMBER: ___ ___ – ___ ___ – ___ ___

ERROR CODE: ____________________  PROBLEM TITLE: ____________________

FRB MEMBER: ____________________  ASSIGNEE: ____________________

TYPE OF PROBLEM: F/A □ O □  CURRENT FIX DATE/SW REV. ____________________

COMMON: □  UNIQUE: □  CATEGORY: WORK: □  PROC: □  DESIGN: □

MODULE CODE: ___ ___

INSUFFICIENT INFO: □ DATE: _________  SUFFICIENT INFO: □ DATE: _________

COMMENTS: ____________________

□ DEFINITION OF PROBLEM: ____________________

______________________________  DEFINED: □ DATE ____________________

□ CONTAINMENT: ____________________

______________________________  CONTAINED: □ DATE ____________________

□ ROOT CAUSE: ____________________

□ CORRECTIVE ACTION: ____________________

______________________________

VERIFICATION: ____________________  RETIRED/CLOSED: □ DATE: _________

DOCUMENTATION: □ ECN: _________  □ FRI  _________  □ OTHER: _________

______________________________

REVISION: A: □ B: □  C: □  D: □  DATE: _________

______________________________

ENTERED IN DB: _________

*** IF MORE SPACE IS NEEDED, PLEASE ATTACH SEPARATE PIECE OF PAPER ***

Figure 10  PAR Form
Failure Review Board

The FRB reviews failure trends, facilitates and manages the failure analysis, and participates in developing and implementing the resulting corrective actions. To do these jobs properly, the FRB must be empowered with the authority to require investigations, analyses, and corrective actions by other organizations. The FRB has much in common with the techniques of quality circles; they are self-managed teams directed to improve methods under their control (see Figure 11).

![FRB Process Diagram]

Figure 11  FRB Process
Figure 11, continued  FRB Process

- Enter into DBMS & write PARs
  - After suitable period if additional events have not recurred, retire problem
- Sufficient information?
  - Yes
  - Establish containment, RCA plan/schedule
  - FRB assigns problem owner
  - Design problems
  - Software problems
  - Mfg. issues
  - Operator/field services
  - Others
  - Establish containment, RCA plan/schedule
- FRB reviews & ensures reliable solution
- RCA complete
- Begin ECN Process
- Update DBMS
- Distribute updates to FRB
- Enter into DBMS & write PARs
  - After suitable period if additional events have not recurred, retire problem
- Sufficient information?
  - No
  - FRB assigns problem owner
  - Design problems
  - Software problems
  - Mfg. issues
  - Operator/field services
  - Others
  - Establish containment, RCA plan/schedule
  - FRB reviews & ensures reliable solution
  - RCA complete
  - Begin ECN Process
  - Update DBMS
The makeup of the FRB and the scope of authority for each member should be identified in the FRACAS procedures. The FRB is typically most effective when it is staffed corporate-wide, with all functional and operational disciplines within the supplier organization participating. The user may be represented also. Members should be chosen by function or activity to let the composition remain dynamic enough to accommodate personnel changes within specific activities.

The FRB composition may be product specific, if desired. For example, a supplier’s different products may be complex and therefore require expert knowledge. In this scenario, though, overall priorities must be managed carefully, since the FRB may require the same expert resources for root cause analysis.

Generally, the following organizations are represented on the FRB:

<table>
<thead>
<tr>
<th>Reliability</th>
<th>Quality Assurance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Field Service</td>
<td>Manufacturing</td>
</tr>
<tr>
<td>Statistical Methods</td>
<td>Marketing</td>
</tr>
<tr>
<td>Systems Engineering</td>
<td>Test</td>
</tr>
<tr>
<td>Design Engineering*</td>
<td></td>
</tr>
</tbody>
</table>

The FRB should be headed by the reliability manager. One of the main functions of this reliability manager is to establish an effective FRACAS organization. The reliability manager must establish procedures, facilitate periodic reviews, and coach the FRB members. Other responsibilities include:

- Assign action items with ownership of problem/solution (who/what/when)
- Allocate problems to appropriate functional department for corrective action

* Hardware, software, process, and/or materials design, depending on the type of system being analyzed.
- Conduct trend analysis and inform management on the types and frequency of observed failures
- Keep track of failure modes and criticality
- Issue periodic reports regarding product performance (for example, MTBF, MTTR, productivity analysis) and areas needing improvement
- Ensure problem resolution
- Continually review the FRB process and customize it to fit specific product applications

The FRB operates most efficiently with at least two recommended support persons. A reliability coordinator can procure failed items for root cause analysis, ensure that the event reports submitted to the FRB contain adequate information, stay in contact with internal and external service organizations to ensure clarity on service reports, and track and facilitate RCA for corrective action implementation.

A database support person should be responsible for providing periodic reports and for maintaining the database (importing/exporting data, backing up files, archiving old records). This will ensure that data available from RCA and corrective action is kept current in the database.

Responsibilities of the FRB members include the following:

- Prepare a plan of action with schedule
- Review and analyze failure reports from in-house tests and field service reports
- Identify failure modes to the root cause
Furthermore, each FRB member should

- Have the authority to speak for and accept corrective action responsibility for an organization
- Have thorough knowledge of front-end systems
- Have the authority to ensure proper problem resolution
- Have the authority to implement RCA
- Actively participate

The FRB should meet on a regular basis to review the reported problems. The frequency of these meetings will depend on the number of problems being addressed or on the volume of daily event reports. Problem-solving skills and effective meeting techniques can ensure that meetings are productive. If you need assistance in these areas, SEMATECH offers short courses\(^3\) that are affordable and can be customized for your application.

The top problems, based on pareto analyses (statistical methods) derived from the database, should be used to assign priorities and to allocate resources.

The reporting relationship of the FRB with other FRACAS functions is shown in Figure 12.
Root Cause Analysis

In failure analysis, reported failures are evaluated and analyzed to determine the root cause. In RCA, the causes themselves are analyzed and the results and conclusions are documented. Any investigative or analytical method can be used, including the following:

- Brainstorming
- Histogram
- Flow chart
- Force field analysis

Figure 12 FRACAS Functions Responsibilities
- Pareto analysis
- Nominal group technique
- FMEA
- Trend analysis
- Fault tree analysis
- Cause and effect diagram (fishbone)

These tools are directly associated with the analysis and problem-solving process. Advanced methods, such as statistical design of experiments, can also be used to assist the FRB. The Canada and Webster divisions of Xerox Corporation extensively use design of experiments as their roadmap for problem solving; they find this method helps their FRBs make unbiased decisions.

Any resulting corrective action should be monitored to ensure that causes are eliminated without introducing new problems.

**Failed Parts Procurement**

A failed parts procurement process should be established to assist the FRB in failure analysis. Failed parts may originate from field, factory, or company stock, or they may become defective during shipment. Each of these scenarios requires different action. The procedure should clearly define the process, including the timeline and responsibilities for at least the following tasks:

- Shipment of failed parts from field to factory
- Procurement of failed parts from various factory locations

---

4. Failure Mode and Effects Analysis. More information is available through the Failure Mode and Effects Analysis (FMEA): A Guide for Continuous Improvement for the Semiconductor Industry, which is available from SEMATECH as technology transfer #92020963A-ENG.
5. Analysis and problems solving tools are described in the Partnering for Total Quality: Total Quality Toolkit, Vol. 6, which is available from SEMATECH as technology transfer #90060279A-GEN.
6. A fuller description of the Xerox process can be found in Proceedings of Workshop on Accelerated Testing of Semiconductor Manufacturing Equipment, which is available from SEMATECH as technology transfer #91050549A-WS.
- Failed part reports from each location
- Communications between material disposition control and reliability engineering
- Sub-tier supplier failure analysis
- Procurement of sub-tier supplier PAR and implementation of corrective action
- Disposition of failed part after RCA is completed

**CORRECTIVE ACTION**

When the cause of a failure has been determined, a corrective action plan should be developed, documented, and implemented to eliminate or reduce recurrences of the failure. The implementation plan should be approved by a user representative, and appropriate change control procedures should be followed. The quality assurance organization should create and perform incoming test procedures and inspection of all redesigned hardware and software. SEMATECH has developed an Equipment Change Control System\(^7\) that you may find useful (see Figure 13).

To minimize the possibility of an unmanageable backlog of open failures, all open reports, analyses, and corrective action suspension dates should be reviewed to ensure closure. A failure report is closed out when the corrective action is implemented and verified or when the rationale is documented (and approved by the FRB) for any instances that are being closed without corrective action.

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7. Refer to *Equipment Change Control: A Guide for Customer Satisfaction*, which is available from SEMATECH as technology transfer #9011448A–GEN.
An effective change control system incorporates the following characteristics.

- An established equipment baseline exists with current and accurate
  - Design specs and drawings
  - Manufacturing process instructions
  - Equipment purchase specifications

- All changes are identified by using
  - The equipment baseline
  - Raw materials supplier change control
  - Procurement methods and procedures
  - Incoming quality control

- All changes are recorded, forecasted, and tracked in an equipment
  change database incorporating
  - Name of change
  - Change description
  - Reason for change
  - All equipment affected
  - Retrofit required/optional
  - Concerns/risks

- Proper investigation of all changes is performed by a Change Evaluation
  Committee made of experts from
  - Equipment manufacturing
  - Equipment design
  - Software design
  - Quality and reliability
  - Service and maintenance
  - Environment, health, and safety
  - Procurement
  - Process engineering
  - Marketing

- A Change Evaluation Committee is chartered to
  - Validate change benefits using statistical rigor as required
  - Specify change qualification and implementation plans
  - Manage and drive change implementation schedules
  - Update equipment specifications and drawings
  - Maintain the equipment change database
  - Communicate with the customer

In summary, a change control system is functioning well when it systemati-
cally provides

- Identification of all changes relative to an equipment baseline
- Recording, forecasting, and tracking of all changes
- Proper investigation of all changes
- Communication to the customer

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Figure 13  Equipment Change Control Characteristics
The FRACAS Database Management System (DBMS) facilitates failure reporting to establish a historical database of events, causes, failure analyses, and corrective actions. This pool of knowledge should minimize the recurrence of particular failure causes. This DBMS is not product specific. It is available from SEMATECH\(^8\), or you can implement the same concept in whatever software you choose. However, the database should have the same software requirements and characteristics at both the supplier’s and user’s sites.

This section describes characteristics of the SEMATECH product. If you are creating your own implementation, you should develop the following aspects. Sections follow that describe each table more fully.

- Configuration Table
- Events Table
- Problems Table
- Reports

\(^8\) Available from SEMATECH Total Quality division. Software transfer and accompanying documentation in press.
**Configuration Table**

The configuration table accepts information about each machine or software instance. There are six standard fields, as shown in the following table. In the SEMATECH FRACAS database, there are seven optional fields that can be customized for your purposes, such as customer identification number, software revision, process type, engineering change number, site location, etc.

**TABLE 2  STANDARD FIELDS OF THE CONFIGURATION TABLE**

<table>
<thead>
<tr>
<th>Field Name</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serial Number</td>
<td>User input.</td>
</tr>
<tr>
<td>Site</td>
<td>Enter the location of the machine.</td>
</tr>
<tr>
<td>System</td>
<td>User input.</td>
</tr>
<tr>
<td>Phase</td>
<td>User input. Enter the current Life Cycle Phase of the machine. Required field.</td>
</tr>
<tr>
<td>Phase Date</td>
<td>User input. Enter the date the machine went into the current Life Cycle Phase.</td>
</tr>
<tr>
<td>Weekly Operational</td>
<td>User input. Enter the number of hours the machine is scheduled to be in operation.</td>
</tr>
</tbody>
</table>
Events Table

The primary purpose of the FRACAS application is to record downtime events for a particular system. The fields of the Events Table are described in Table 3. To understand these fields fully, you need to understand the relationship between the problem fields (System, Subsystem, Assembly, Subassembly, and/or Sub-subassembly) and the reliability code.

The reliability code field contains one or more of the problem field names. Data (if any) from the problem fields automatically helps you select or create the reliability code. Once you select a code, associated fields are updated automatically.

TABLE 3 EVENTS TABLE FIELD DESCRIPTIONS

<table>
<thead>
<tr>
<th>Field Name</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serial Number</td>
<td>User input. The serial number of machine having event.</td>
</tr>
<tr>
<td>Date</td>
<td>User input. The date the event occurred. Enter in a MM/DD/YY format.</td>
</tr>
<tr>
<td>Time</td>
<td>User input. The starting time of the event. Entered in 24-hour time (HH:MM:SS). Seconds are not required.</td>
</tr>
<tr>
<td>Duration</td>
<td>User input. The duration of the event. Entered in a HH:MM:SS format. Seconds are not required.</td>
</tr>
<tr>
<td>System</td>
<td>Automatic input based upon the serial number.</td>
</tr>
<tr>
<td>Subsystem</td>
<td>User input. The optional subsystem code (if the part is at a subsystem level) involved in the event. Related to the System codes.</td>
</tr>
<tr>
<td>Assembly</td>
<td>User input. The optional assembly code (if the part is at an assembly level) involved in the event. Related to the selected System and Subsystem codes.</td>
</tr>
<tr>
<td>Operator</td>
<td>User input. Name of the operator when the event occurred.</td>
</tr>
</tbody>
</table>

continued on next page
<table>
<thead>
<tr>
<th>Field Name</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSE Name</td>
<td>User input. Name of field service engineer who responded to the event.</td>
</tr>
<tr>
<td>FSR#</td>
<td>User input. The field service report number.</td>
</tr>
<tr>
<td>Subassembly</td>
<td>User input. The optional subassembly code (if the part is at a subassembly level) involved in the event. Related to the selected System, Subsystem, and Assembly codes.</td>
</tr>
<tr>
<td>Sub-subassembly</td>
<td>User input. The optional sub-subassembly code (if the part is at a sub-subassembly level) involved in the event. Related to the selected System, Subsystem, Assembly, and Subassembly codes.</td>
</tr>
<tr>
<td>Reliability Code</td>
<td>User input. The problem reliability code. This is validated against current reliability codes. A new one may be added by selecting “New” from the pop-up list box.</td>
</tr>
<tr>
<td>Life Cycle Phase</td>
<td>The current phase of the system. Defaults to current phase from Configuration file.</td>
</tr>
<tr>
<td>Down Time Category</td>
<td>User input. Downtime code. Either scheduled or unscheduled. Scheduled downtime should be used for holidays, weekends, or any other time when the machine is not scheduled to be in operation.</td>
</tr>
<tr>
<td>Part Number</td>
<td>User input. The machine part number.</td>
</tr>
<tr>
<td>Relevant Failure</td>
<td>User input. Whether the failure was relevant or not. Y for yes; N for no.</td>
</tr>
<tr>
<td>PM</td>
<td>User input. Whether the downtime was due to preventive maintenance. Y for yes; N for no.</td>
</tr>
<tr>
<td>Problem</td>
<td>User input. The description of the problem. This can be any number of characters.</td>
</tr>
<tr>
<td>Repair Action</td>
<td>User input. The description of the repair action. This can be any number of characters.</td>
</tr>
</tbody>
</table>
Problems Table

In the SEMATECH implementation, the Problem Record allows you to view, modify, or add records for problems. The fields of the Problems Table are described in Table 4.

### Table 4 Problems Field Descriptions

<table>
<thead>
<tr>
<th>Field Name</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>System</td>
<td>User input from Events screen. If adding from Events screen, automatic input based upon the serial number.</td>
</tr>
<tr>
<td>Subsystem</td>
<td>User input. Optional subsystem code (if part is subsystem level) involved in event. Related to System code. See also Reliability Code.</td>
</tr>
<tr>
<td>Assembly</td>
<td>User input. Optional assembly code (if part is at assembly level) involved in event. Related to selected System and Subsystem codes. See also Reliability Code.</td>
</tr>
<tr>
<td>Subassembly</td>
<td>User input. Optional subassembly code (if part is at subassembly level) involved in event. Related to selected System, Subsystem, and Assembly codes. See also Reliability Code.</td>
</tr>
<tr>
<td>Sub-subassembly</td>
<td>User input. Optional sub-subassembly code (if part is at sub-subassembly level) involved in event. Related to selected System, Subsystem, Assembly, and Subassembly codes. See also Reliability Code.</td>
</tr>
<tr>
<td>Reliability Code</td>
<td>Automatically generated from the above fields. A sequence number will be added to the code.</td>
</tr>
<tr>
<td>Title/Description</td>
<td>User input. Simple title or description of event.</td>
</tr>
<tr>
<td>Initial Status</td>
<td>User input. Required. Status of problem when first reported. If problem is being added from Event screen, date of event is entered automatically as initial status date.</td>
</tr>
<tr>
<td>Fault Category</td>
<td>User input. Event category.</td>
</tr>
<tr>
<td>Error Codes</td>
<td>User input. Error code for selected tool.</td>
</tr>
<tr>
<td>Assignee</td>
<td>User input. Staff member who assigned this problem.</td>
</tr>
<tr>
<td>Insufficient Information</td>
<td>Date when insufficient information status was declared.</td>
</tr>
</tbody>
</table>
### Table 4: Problems Field Descriptions, Continued

<table>
<thead>
<tr>
<th>Field Name</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sufficient Information</td>
<td>Date when sufficient information status was declared.</td>
</tr>
<tr>
<td>Defined</td>
<td>Date when defined information status was declared.</td>
</tr>
<tr>
<td>Contained</td>
<td>Date when contained information status was declared.</td>
</tr>
<tr>
<td>Retired</td>
<td>Date when retired information status was declared.</td>
</tr>
<tr>
<td>Root Cause Solution – Original</td>
<td>Original (planned) date for fixing problem.</td>
</tr>
<tr>
<td>Root Cause Solution – Current</td>
<td>Current date for fixing problem.</td>
</tr>
<tr>
<td>Complete</td>
<td>Completion date for fixing problem.</td>
</tr>
<tr>
<td>RCS Documentation</td>
<td>User input. Points to documentation of fix; for example, Engineering Change Numbers (ECNs).</td>
</tr>
<tr>
<td>Comments</td>
<td>User input. Memo area for problem comments. Can be any number of characters.</td>
</tr>
</tbody>
</table>

### Report Types

Your reporting mechanism for this database should be flexible enough to accommodate various customer types and their needs. For example, you need to be able to pull reports for an FRB analysis as well as for the users, sub-tier suppliers, and those who are addressing corrective actions. At a minimum, the following reports should be available:

- Trend charts to show a number of events during a time period.
- Event history to show a list of events for during a time period.
- Problem history to show a list of problems during a time period.
- Reliability statistics to demonstrate the system performance in terms of MTTR, MTBF, mean time to failure (MTTF), availability, etc.
GLOSSARY

Corrective Action  A documented change to design, process, procedure, or material that is implemented and proven to correct the root cause of a failure or design deficiency. Simple part replacement with a like item is not considered corrective action.

Equipment Life Cycle  The sequence of phases or events constituting total product existence. The equipment life cycle is divided into the following six phases:
  - Concept and feasibility
  - Design
  - Prototype (alpha-site)
  - Pilot production (beta-site)
  - Production and operation
  - Phase-out

Error  Human action that results in a fault (term is usually reserved for software).

Error Code  Numbers and/or letters reported by the equipment’s software that represent an error type; code helps determine where the fault may have originated.

Failure  An event in which an item does not perform its required function within the specified limits under specified conditions.

Failure Analysis  A determination of failure cause made by use of logical reasoning from examination of data, symptoms, available physical evidence, and laboratory results.
**Failure Cause** The circumstance that induces or activates a failure. Examples of a failure cause are defective soldering, design weakness, assembly techniques, and software error.

**Failure Mode** The consequence of the failure mechanism (the physical, chemical, electrical, thermal, or process event that results in failure) through which the failure occurs; for example, short, open, fracture, excessive wear.

**Failure Review Board (FRB)** A group consisting of representatives from appropriate organizations with the level of responsibility and authority to assure that failure causes are identified and corrective actions are effected.

**Fault** Immediate cause of failure. A manifestation of an error in software (bug), if encountered, may cause a failure.

**Fault Code** Type of failure mechanism categorized to assist engineering in determining where the fault may have originated. For example, a fault may be categorized as electrical, software, mechanical, facilities, human, etc.

**FMEA** Failure Modes and Effects Analysis. An analytically derived identification of the conceivable equipment failure modes and the potential adverse effects of those modes on the system and mission.

**FRACAS** Failure Reporting, Analysis, and Corrective Action System. A closed-loop feedback path by which failures of both hardware and software data are collected, recorded, analyzed, and corrected.
Laboratory Analysis  The determination of a failure mechanism using destructive and nondestructive laboratory techniques such as X-ray, dissection, spectrographic analysis, or microphotography.

Mean Time to Repair (MTTR)  Measure of maintainability. The sum of corrective maintenance times at any specific level of repair, divided by the total number of failures within an item repaired at that level, during a particular interval under stated conditions.

Mean Time Between Failures (MTBF)  Measure of reliability for repairable items. The mean number of life units during which all parts of the item perform within their specified limits, during a particular measurement interval under stated conditions.

Mean Time to Failure (MTTF)  Measure of reliability for nonrepairable items. The total number of life units of an item divided by the total number of failures within that population, during a particular measurement interval under stated conditions.

Related Event  A failure that did not cause downtime. A related event or failure is discovered while investigating a failure that caused downtime. A related event could also be a repair or replacement during downtime caused by the main event.

Relevant Failure  Equipment failure that can be expected to occur in field service.

Reliability  The duration or probability of failure-free performance under stated conditions.

Reliability Code  A functional traceable description of the relationships that exists in the equipment. Codes are
derived in a tree fashion from top to bottom as System, Subsystem, Assembly, Subassembly, and Sub-subassembly (typically lowest replaceable component). Codes vary in size; however, a maximum three alphanumeric descriptor per level is recommended. For example, AH-STK-ROT-XAX-MOT for Automated Handler-Stock-Robot-X_Axis-Motor.

**Repair or Corrective Maintenance** All actions performed as a result of failure to restore an item to a specified condition. These can include localization, isolation, disassembly, interchange, reassembly, alignment, and checkout.

**Root Cause** The reason for the primary and most fundamental failures, faults, or errors that have induced other failures and for which effective permanent corrective action can be implemented.

**Uptime** Time when equipment is in a condition to perform its intended function. It does not include any portion of scheduled downtime and nonscheduled time.

**Glossary References:**


# FRACAS Functions and Responsibilities

<table>
<thead>
<tr>
<th>Responsibilities</th>
<th>Function</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Failure</td>
<td>Operators</td>
<td>Identify problems. Call for maintenance. Annotate incident.</td>
</tr>
<tr>
<td>Failure Report</td>
<td>Maintenance</td>
<td>Generate report with supporting data (time, place, equipment, etc.).</td>
</tr>
<tr>
<td>Data Logging</td>
<td>Reliability</td>
<td>Log all failure reports. Validate failures and forms. Classify failures (inherent, induced, false alarm, etc.).</td>
</tr>
<tr>
<td>Failure Analysis</td>
<td>Reliability and/or Problem Owners</td>
<td>Review operating procedures for error. Procure failed parts. Decide which parts will be destructively analyzed. Perform failure analysis to determine root cause.</td>
</tr>
<tr>
<td></td>
<td>Quality</td>
<td>Inspect incoming test data for item.</td>
</tr>
<tr>
<td>Corrective Action</td>
<td>Design</td>
<td>Redesign hardware/software, if necessary.</td>
</tr>
<tr>
<td></td>
<td>Sub-tier Supplier</td>
<td>Prepare &amp; provide new part or test procedure, if necessary.</td>
</tr>
<tr>
<td></td>
<td>Quality</td>
<td>Evaluate incoming test procedures. Inspect redesigned hardware/software.</td>
</tr>
<tr>
<td>Post-data Review</td>
<td>Reliability</td>
<td>Close loop by collecting and evaluating post-test data for recurrence of failure.</td>
</tr>
</tbody>
</table>