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**Results of the  
IEC 61508  
Functional Safety Management  
Assessment**

Customer:

**PR electronics**

Rønde,  
Denmark

Contract No.: 0709-02C

Report No.: 0709-02C R004 Assessment

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Peter Müller

## Management summary

The Functional Safety Assessment of the PR electronics, performed by *exida* Certification S.A. consisted of the following activities:

- *exida* Certification S.A. assessed the setup of the development process used by PR electronics for development projects against the relevant requirements of IEC 61508 parts 1 to 3.

Subject to this assessment were the Functional Safety Planning activities, the tailoring of the Verification and Validation activities and the realization of the technical safety aspects using an actual safety development project as example.

- *exida* Certification S.A. audited the development process by a detailed development audit which investigated the compliance with IEC 61508 of the processes, procedures and techniques as implemented for the PR electronics safety products development. The investigation was executed using subsets of the IEC 61508 requirements tailored to the work scope of the development team.



Some areas for improvement were identified which are generally required to formally show the compliance to IEC 61508. However, because of the size of the project (limited number of people) and the low complexity / limited size of the product, PR electronics was able to demonstrate that the *objectives of the related areas have been successfully met*.

The result of the Functional Safety Assessment can be summarized by the following statements:

**The assessment has shown that the Hardware and Software development processes in general, complies with the relevant safety requirements for design, implementation and verification for IEC 61508 SIL2.**

**The audited PR electronics development process tailored and implemented by the safety related Hardware and Software development projects, complies with the relevant functional safety management requirements of IEC 61508 SIL2.**

**This means that products, with Hardware and Software, developed according to the functional safety management system are capable for use in SIL2 applications, related to the measures to avoid systematic faults during the product development.**

	
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<i>exida</i> Certification S.A. Peter Müller	pr 0709-02c assessment report fsm certificate v1r1.doc, 28 November 2008 Page 2 of 19

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## 1 Purpose and Scope

This document describes the results of the IEC 61508 Functional Safety Management Assessment of the product development processes according to the safety lifecycle phase 9 of IEC 61508-1. The purpose of the assessment was to investigate the compliance of

- the product development processes, procedures and techniques as implemented for the safety-related deliveries with the managerial IEC 61508-1, -2 and -3 requirements for SIL2.

It was not the purpose of the assessment / audits to investigate Company quality management system versus ISO 9001 and ISO 9000-3 respectively.

The assessment has been carried out based on the quality procedures and scope definitions of *exida* Certification S.A..

## 2 Description of the Functional Safety Management System

The functional management system is implemented by the use of the functional safety management plan and the related planning documents, which describes the safety related processes in detail. The functional safety management plan shows the implementation of a safety life cycle model which adopts the V-model as described in IEC 61508.

The related planning documents are mainly the configuration management plan, the verification and validation plan and a set of guidelines.

Evidence for the fulfilment of the detailed requirements have been collected in a Safety Justification report, which was subject to the assessment.

## 3 Project management

### 3.1 Assessment of the development process

The development audit was closely driven by requirements subsets filtered from the IEC 61508 content of the *exida* SafetyCaseDB database. That means that the Functional Safety Management related requirements were grouped together according their related objectives. The detailed answers to the requirements, i.e. the justification report, were subject to the assessment. This assessment of the justification report was supplemented by the prior review of documents.

The assessment was planned by *exida* Certification S.A. and agreed with PR electronics.

The following IEC 61508 objectives were subject to detailed auditing at PR electronics:

- FSM planning, including
  - Safety Life Cycle definition
  - Scope of the FSM activities
  - Documentation
  - Activities and Responsibilities (Training and competence)
  - Configuration management
  - Tools and languages
- Safety Requirement Specification
- Change and modification management
- Software architecture design process, techniques and documentation
- Hardware architecture design - process, techniques and documentation
- Hardware and system related V&V activities including documentation, verification
  - Integration and fault insertion test strategy
- Software and system related V&V activities including documentation, verification

- System Validation including hardware and software validation
- Hardware-related operation, installation and maintenance requirements

The project teams, not individuals were audited.

The development audit was executed in several steps.

A pre-assessment of the safety relevant documents has been performed off site.

The audits have been done in Rønne, Denmark at 2007.12.17 – 19, 2008.02.19 – 22 and 2008.04.21 -23.

### 3.2 Roles of the parties involved

#### PR electronics

Represents the designer of the safety related products and the investigated organization. The following teams / responsible persons were audited:

- |                               |                            |
|-------------------------------|----------------------------|
| • Project & Safety Management | Hans Jørgen Eriksen        |
| • Hardware development        | Hans Jørgen Eriksen        |
| • Software development        | Flemming Svanholm Sørensen |
| • Test leader                 | Kaj Harbo                  |

#### *exida* Certification S.A.

Set up and structure of the assessment and audit process, extracted the requirements for the assessment and audit from the IEC 61508 standard and guided through the audit.

The activities were done by *exida* Certification S.A. as an independent organization. The assessment was performed by Peter Müller, who was not involved in the execution of the audited activities.

## 4 Results of the Functional Safety Assessment

exida Certification S.A. assessed the development process used by PR electronics for development projects against the objectives of IEC 61508 parts 1 to 3. The results of the pre-assessment are documented in [R1].

All objectives have been successfully considered in the PR electronics development processes for their safety related product development.

exida Certification S.A. assessed the safety justification report prepared by PR electronics, a set of documents, against the functional safety management requirements of IEC 61508. This was done by a pre-review of the completeness of the related requirements and then a spot inspection of certain requirements, before the development audit.

The safety justification report demonstrated the fulfillment of the functional safety management requirements of IEC 61508-1 to 3.

The detailed development audit (see [R1]) investigated the compliance with IEC 61508 of the processes, procedures and techniques as implemented for the PR electronics functional safety management system.

The investigation was executed using subsets of the IEC 61508 requirements tailored to the work scope of the development team. The result of the assessment can be summarized by the following observations:

**The assessment has shown that the Hardware and Software development processes in general, complies with the relevant functional safety management requirements for design, implementation and verification for IEC 61508 SIL2.**

**The audited development process tailored and implemented for safety related Hardware and Software development projects, complies with the relevant functional safety management requirements of IEC 61508 SIL2.**

**This means that products, with Hardware and Software, developed according to the functional safety management system are capable for use in SIL2 applications, related to the measures to avoid systematic faults during the product development.**

Some areas for improvement were nevertheless identified. The recommended improvements given below (Rec: x) are generally required to formally show the compliance to IEC 61508. However, because of the size of the project (limited number of people) and the low complexity / limited size of the products, PR electronics was able to demonstrate that the *objectives of the related areas have been successfully met*. More details can be found in the chapters below.

### 4.1 Functional Safety Management

#### Objectives of the Functional Safety Management

The main objectives of the related IEC 61508 requirements are to:

- Structure, in a systematic manner, the phases in the overall safety lifecycle that shall be considered in order to achieve the required functional safety of the E/E/PE safety-related systems.

- Structure, in a systematic manner, the phases in the E/E/PES safety lifecycle that shall be considered in order to achieve the required functional safety of the E/E/PE safety-related systems.
- Specify the management and technical activities during the overall, E/E/PES and software safety lifecycle phases which are necessary for the achievement of the required functional safety of the E/E/PE safety-related systems.
- Specify the responsibilities of the persons, departments and organizations responsible for each overall, E/E/PES and software safety lifecycle phase or for activities within each phase.
- Specify the necessary information to be documented in order that the management of functional safety, verification and the functional safety assessment activities can be effectively performed.
- Document all information relevant to the functional safety of the E/E/PE safety-related systems throughout the E/E/PES safety lifecycle.
- Document key information relevant to the functional safety of the E/E/PE safety-related systems throughout the overall safety lifecycle.
- Specify the necessary information to be documented in order that all phases of the overall, E/E/PES and software safety lifecycles can be effectively performed.
- Select a suitable set of tools, for the required safety integrity level, over the whole safety lifecycle which assists verification, validation, assessment and modification.

#### 4.1.1 Safety Life Cycle

The development process is well structured and described in the 9000 FSM Plan. It describes all relevant phases for development, integration, verification, validation and modification. The related activities including inputs and outputs assumed for each phase are described.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

#### 4.1.2 FSM planning

The 9000 FSM Plan defines for the different work items the required input documents, guidelines and templates. The phases are specified in the 9000 FSM Plan and the 9000 V&V plan. All major activities related to specification, verification and validation are planned in the 9000 FSM Plan. The different roles and responsibilities of people are defined in the 9000 RACI chart. The modification procedure after product release is part of this document.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

#### 4.1.3 Documentation

All V&V specifications, and reports are kept under version control together with the associated design and product documents.

The test specification templates describes precisely how to document the validation and integration tests, their specifications, their execution and the results. The templates enables the re-execution of tests by requiring the relevant information.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

#### 4.1.4 Training and competence recording

The FSM Plan have been specified, reviewed and approved by the responsible people for the specified activities of the project.

The responsibility for the documents are tracked in the RACI chart.

The FSM plan requires to collect the evidence documentation regarding the competence of the involved parties in the project. This is documented in the competence matrix document.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

#### 4.1.5 Configuration Management

All work products are part of a Visual Source Safe based version management system.

The HW and SW modules building the subsystem can be identified by a naming / numbering convention as described in the Q-system (KMH). The project documents are listed / defined in the RACI-chart together with their version and revision.

The connection between these named items, their version / revision and (internal) releases (baselines, labels, builds, etc) can be generated out of the SourceSafe database. In the Correction sheet for each product the connection between the firmware and hardware version is listed.

There is a set of master copy(ies) / Baselines available that contains all work products that were used as an argument for demonstrating safety integrity of a certain version.

Which versions of a work product was part of which test run is documented in the respective test reports.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

#### **4.1.6 Tools (and languages)**

The 9000 FSM Plan and the "9000 Confidence from Use of Software tools" lists the selected set of tools and argues for their suitability.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

## **4.2 Safety Requirement Specification**

### **Objectives of the Safety Requirement Specification**

The main objectives of the related IEC 61508 requirements are to:

- Specify the requirements for each E/E/PE safety-related system, in terms of the required safety functions and the required safety integrity, in order to achieve the required functional safety.

#### **4.2.1 Safety Requirement Specification and traceability into design**

The FSM plan requires the SRS to be developed before any other design and development activity as input for the architecture design of the system / product. For the System9000 project, the final SRS and Safety concept iterations was developed partly in parallel with the development activities.

For each product (sometimes product pairs) one SRS is existing covering all technical safety requirements, both for system and SW, with a clear identification of safety and non-safety related requirements.

The structure and consistency of the SRS is achieved through use of a template back-end, which is based on the IEC 61508 standard.

During the architectural system and software design, the SRS is reviewed by designers for completeness and understandability. The target of the review is always to detect inconsistencies and incompatibilities of the requirements.

The safety concept contains references to the requirements in the SRS. This enables a verification of that the architecture is addressing all applicable requirements in the SRS.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

### **4.3 Change and modification management**

#### **Objectives of change and modification management**

The main objectives of the related IEC 61508 requirements are to:

- Ensure that the required safety integrity is maintained after corrections, enhancements or adaptations to the E/E/PE safety-related systems.

#### **4.3.1 Change and modification procedure**

The FSM plan includes a section which describes the modification process. This includes:

- (1) Change request either by a fault found during integration / validation, functional enhancement request or by a (field) failure investigation;
- (2) Impact analysis of the proposed change to the PES itself;
- (3) Specification of the change;
- (4) The impact analysis determines the appropriate re-entry point of the safety life cycle;
- (5) Implement the specified change;
- (6) Re-verification of changed modules and affected modules.
- (7) Re-validation of affected requirements and regression tests;
- (8) Procedures and decision to inform customers upon detection of safety critical faults in released products, these are part of the normal company quality procedures.
- (9) The modification process shall be used starting with formal integration test.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

### **4.4 Software Design**

#### **Objectives of software design**

The main objectives of the related IEC 61508 requirements are to:

- Create a software architecture that fulfils the specified requirements for software safety with respect to the required safety integrity level.
- Review and evaluate the requirements placed on the software by the hardware architecture of the E/E/PE safety-related system, including the significance of E/E/PE hardware/software interactions for safety of the equipment under control.
- Design and implement software that fulfils the specified requirements for software safety with respect to the required safety integrity level, which is analyzable and verifiable, and which is capable of being safely modified.

#### **Objectives of tools and languages**

The main objectives of the related IEC 61508 requirements are to:

- Select a suitable set of tools, including languages and compilers, for the required safety integrity level, over the whole safety lifecycle of the software which assists verification, validation, assessment and modification.

#### 4.4.1 Software architecture design

The design is described by the used UML model in combination with the detailed design description.

The UML subset used addresses the following objectives:

1. Static design - Deployment and Component diagrams
2. Dynamic behavior - State transition diagrams;  
- Sequence diagrams or Object interaction diagrams.
3. Link to the source code - Class diagrams with each .c/.h. pair modeled by a class;

The use of UML supports the need for transparency, abstraction and modularity as required by the "9000 Safety Concept Design using UML".

The use of this design method is supported by the software tool Enterprise Architect which is used for safety related projects.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

#### 4.4.2 Tools and languages

For the System 9000 the compiler vendor provides a statement of the compliance with well accepted test suites like Plum Hall (ANSI C). This is documented in the "Confidence from use of software tools" document.

The "9000 Style Guide for Firmware Coding" describes the coding standard for this project. It is based on the MISRA coding standard together with some PR electronics defined stricter rules. The source is checked by PC-Lint, a static code analysis tools together with the applied MISRA rules. Rules that cannot be automatically checked are part of the checklist for manual source code review.

The "9000 Style Guide for Firmware Coding" additionally describes the "style guides" for the source code files / documentation regarding description, inputs and output. Also naming conventions, information requirements and layout of the files are described here

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

### 4.5 Hardware Design

#### Objectives of hardware design

The main objectives of the related IEC 61508 requirements are to:

- Create E/E/PE safety-related systems conforming to the specification for the E/E/PES safety requirements (comprising the specification for the E/E/PES safety functions requirements and the specification for the E/E/PES safety integrity requirements).

- Ensure that the design and implementation of the E/E/PE safety-related systems meets the specified safety functions and safety integrity requirements.

#### **Objectives of hardware design / probabilistic properties**

The main objectives of the related IEC 61508 requirements are to:

- Ensure that the design and implementation of the E/E/PE safety-related systems meets the specified safety functions and safety integrity requirements.

#### **4.5.1 Hardware architecture design**

There is a description of the HW architecture in the safety concept document.

The sub-systems with their HW / SW and SW / SW interactions are specified and documented together with their safety relevance in the Safety Criticality Analysis report (in the System-FMEA) / architecture description.

The HW/HW interactions are described in more detail in the different circuit description documents. This serves both as input to specification of integration tests and as information about which functions and interfaces that can be used by safety functions.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

#### **4.5.2 Hardware Design / Probabilistic properties**

The detailed hardware design is described by Circuit Diagrams, layout drawings and a related parts list. As required by IEC 61508, an FMEDA with probabilistic calculations and the related fault insertion tests will be carried out for the safety related products, as planned by the 9000 FSM plan.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

### **4.6 Verification & Validation**

#### **Objectives of HW related verification & validation activities**

The main objectives of the related IEC 61508 requirements are to:

- Demonstrate, for each phase of the overall, E/E/PES and software safety lifecycles (by review, analysis and/or tests), that the outputs meet in all respects the objectives and requirements specified for the phase.
- Test and evaluate the outputs of a given phase to ensure correctness and consistency with respect to the products and standards provided as input to that phase.
- Integrate and test the E/E/PE safety-related systems.

- Ensure that the design and implementation of the E/E/PE safety-related systems meets the specified safety functions and safety integrity requirements.
- Plan the validation of the safety of the E/E/PE safety-related systems.
- Validate that the E/E/PE safety-related systems meet, in all respects, the requirements for safety in terms of the required safety functions and the safety integrity.

#### **4.6.1 HW related V&V activities**

The V&V Plan specifies the techniques and the project specific tools / test SW which are used in the verification activities for each phase and each product. The criteria are addressed wherever applicable, e.g. for test coverage.

All planned test levels, module-, integration-, fault insertion- and validation-test are specified in accordance to the selected Safety Integrity Level.

All analytical verification activities are described by the combination of FSM plan and V&V Plan.

All validation activities are documented as required by the planning documents. This includes the techniques and methods to be used, e.g. procedural (review) and technical (functional test). The purpose is to show that the system and SW requirements are successfully met.

The selected Requirements Tracking methodology shows traceably the link between safety requirements, validation tests and design. The target is 100% coverage of the safety requirements. The test cases (called test objectives) are reviewed against the validation objectives and the corresponding requirement. The test execution results are reviewed against expected results.

E.g. the 9202 Requirement traceability matrix contains all requirements for the product, both safety and non-safety. This matrix is used to ensure completeness of the test cases versus all requirements.

Each validation test case defines a test objective, test preparation, test steps and expected output including additional acceptance criteria (typically for performance / usability requirements) where applicable.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

#### **4.6.2 SW related V&V activities**

##### **Objectives of SW related verification and validation activities**

The main objectives of the related IEC 61508 requirements are to:

- To the extent required by the safety integrity level, test and evaluate the outputs from a given software safety lifecycle phase to ensure correctness and consistency with respect to the outputs and standards provided as input to that phase.

- Verify that the requirements for software safety (in terms of the required software safety functions and the software safety integrity) have been achieved.
- Integrate the software onto the target programmable electronic hardware. Combine the software and hardware in the safety-related programmable electronics to ensure their compatibility and to meet the requirements of the intended safety integrity level.
- Ensure that the integrated system complies with the specified requirements for software safety at the intended safety integrity level.

The V&V Plan specifies the techniques and the project specific tools / test SW which are used in the verification activities for each phase and each product. The criteria are addressed wherever applicable, e.g. for test coverage.

All planned test levels, module-, integration-, fault insertion- and validation-test are specified in accordance to the selected Safety Integrity Level.

All analytical verification activities are described by the combination of FSM plan and V&V Plan.

The integration test strategy for the integration levels SW-SW and SW-HW are planned and described in the FSM and V&V plan.

The details regarding the tests, test type, test data and expected result / pass-fail criteria are all described in the test specifications (reports).

In the review of the test report, the test results are reviewed against the expected result / pass-fail criteria leading to a conclusion regarding successful completion of test.

The integration test specification uses the safety concept and the UML model together with the interface description defined therein as input documents in order to define the actual integration tests.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

## 4.7 Safety Manual

### Objectives of the Safety Manual

The main objectives of the related IEC 61508 requirements are to:

- Develop procedures to ensure that the required functional safety of the E/E/PE safety-related systems is maintained during operation and maintenance.

#### 4.7.1 Operation, installation and maintenance requirements

The Safety Manual is part of the User Manual and will, for some important information, contain pointers to information in the User manual instead of repeating it.

The Safety Manual of the product documents the following aspects / characteristics in order to enable the end-user to integrate, operate and maintain the "Compliant Item" in his application:

- Limitations of the product and its application / operational environment;
- The achievable highest level of SIL of each sub-system (based on the techniques and measures documented in the safety justification reports);
- Useful lifetime, i.e. components as identified by the FMEDA, where the estimated PF is valid;
- Guidance on recommended periodic (offline) proof test activities / interval for the product;
- Information as provided by the FMEDA:
  - HW fault tolerance;
  - $\Lambda(du)$ ,  $\Lambda(dd)$ ,  $\Lambda(su)$ ,  $\Lambda(sd)$ ,  $\Lambda(\text{no effect, i.e., on dangerous or safe})$
  - safe failure fraction (SFF);
  - diagnostic coverage derived according to IEC61508-2, annex C;
  - diagnostic test interval.
- All system functions and parameters accessible by the user to implement the safety functions;
- User configuration and programming of the safety functions;
- All safety-related interfaces (I/O, communication, HMI) and their performance characteristics;
- All safety-related aspects regarding installation, commissioning, modification and de-commissioning of the product;
- Guidance on operation of the product including assumed organizational measures to protect against operator mistakes;

FMEDA has been chosen as the systematic method to identify failures which are revealed or unrevealed by the cyclic diagnostics. Periodic proof test procedures are developed for any dangerous undetected faults and documented in the Safety Manual.

**Conclusion:** The objectives of the standard are fulfilled by the PR electronics functional safety management system.

Areas of possible improvements have been identified during the assessment. However, these are not assessed to be in contradiction to an overall positive judgment of the subject.

## 5 Reference documents

The services delivered by *exida* Certification S.A. were performed based on the following standards.

- N1 IEC 61508-1:1998 Functional Safety of E/E/PES; General requirements
- N2 IEC 61508-2:2000 Functional Safety of E/E/PES; Hardware requirements
- N3 IEC 61508-3:1998 Functional Safety of E/E/PES; Software requirements

The pre-assessment delivered by *exida* Certification S.A. were performed based on the audit of the following documents.

- D1 Functional Safety Management Plan Project System 9000 V5R0
- D2 Functional Safety Management Justification Report in System 9000 IEC61508 FSM SafetyCaseDB
- D3 9000 Verification & Validation Plan V2R0
- D4 Technical Justification Report in 9202 SafetyCaseDB – Requirements & Solutions V0R21
- D5 Technical Justification Report in 9202 SafetyCaseDB – Validation Objectives V0R21
- D6 9000 Configuration Management Plan
- D7 9202 Safety Requirement Specification V4R0
- D8 9202 Safety Concept V3R0
- D9 9202 System FMEA / Safety Criticality Analysis V1R0

In addition, the following documents were presented by PR electronics during the audit or given to the assessor for review after the audit:

- D10 9000 SRS Review Record
- D11 Requirements Traceability Matrix Review Template
- D12 9000 Baseline Log
- D13 9000 RACI Chart
- D14 9000 Competence of People
- D15 9000 Confidence From-Use of Software Tools V2R0
- D16 Supplier Statements Regarding ANSII Compliance (ZIP file)
- D17 9000 Product History Document
- D18 Quality Handbook - Kvalitets og miljø handbog
- D19 Quality Procedure: Calibration
- D20 9000 Safety Concept Design using UML V1R0

- D21 9000 Style Guide for Firmware Coding V1R0
- D22 9000 Code Review Template
- D23 9000 Integration Test Report Template
- D24 9000 Firmware Design Specification Review Template
- D25 Acceptance Test Report Review Template
- D26 9202 Acceptance Test Report V5R0
- D27 9000 Change Request Template
- D28 920262xx Software Module Test Report
- D29 9202 Software Fault Insertion Test Report V3R0
- D30 9202 Integration Test Report
- D31 9202 Requirement Traceability Matrix
- D32 9202-1-03 Correction Sheet
- D33 9202 Circuit Description Pulse Isolator V4R0
- D34 Safety Manual Pulse Isolator 9202 Version No. V1R0
- D35 92026xxx Firmware Design Specification (UML model)  
(920260xx V7R0; 920261xx V2R0; 920262xx V2R0; 920264xx V3R0)
- D36 System 9000 Relay Endurance Test V0R3
- D37 9202 FMEDA Report V1R0
- D38 9202 Type A FMEDA – Excel Sheets
- D39 9000 LED and Error Indications V4
- D40 9202 Requirement Specification V5R0
- D41 9202 De-rating Analysis for schematics V4R2
- D42 9202 Schematics 9202-1-04-PDF.pdf dated 2008-03-28
- D43 9202 Hardware Fault Insertion Report V3R0
- D44 9202 Hardware Module Test report V6R0

The supporting services delivered by *exida* were documented by the following documents.

- R1 Document Review & Assessment Comments, Version 1, Revision 6, September 2008 Confidential Report.
- R2 Results of the IEC 61508 Functional Safety Management Assessment (this document).
- R3 Minutes of Meeting, 2007.09.4 -7, System 9000 Assessment Preparation
- R4 Assessment Plan, Version 2, Revision 2, February 2008
- R5 Recommendations caused by the IEC 61508 Functional Safety Assessment V1R1, November 2008 Confidential Report.

## 6 Status of the document

### 6.1 Releases

Version History: V0, R1: Development Audit, Initial Report March 19, 2008  
V0, R2 Recommendations moved to [R5], updated according to customers Review, Document numbers updated 02.09.2008,  
V1, R0 Review comments incorporated.  
V1, R1 *exida* Certification logo exchanged, Reference to Safety Manual updated

Author: Peter Müller

Review: V0, R1 Customer  
V0, R2 Rachel van Beurden-Amkreutz

Release status: Released