

# Supplier Quality Requirements

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#### **DOCUMENT HISTORY**

| Issue No | Date       | Brief Description  |
|----------|------------|--|
| 1        | 25/11/2013 | Original document, initial release   |
| 2        | 19/12/2013 | Addition of clause 1.7   |
| 3        | 13/02/2017 | Addition of clause 11, Change of originator job role & update of contents page   |
| 4        | 06/12/2018 | Change of originator job role  |
| 5        | 10/01/2022 | RCCN 344. Format changes in line with current AQWSR documents. Change of creation, approval and sign off personnel. Updated logos. Updates to the following sections: 1, 2, 3, 4, 7, 8, 10, 11. Addition of sections 12, 13, 14. |
| 6        | 29/11/2022 | Changes to the document to capture the requirements of AS9100D clause 8.4.3  |
| 7        | 24/07/2025 | Updates to the formatting of the document and updates against AS9100 requirements.   |



## 1. Scope and Applicability

- This manual applies to all external providers (suppliers, subcontractors, distributors) that deliver products or services impacting AQ Wiring Systems Rockford Ltd.'s quality, safety, and regulatory compliance.
- It includes direct suppliers and sub-tier providers involved in design, manufacturing, testing, special processes, and logistics.
- The requirements are applicable regardless of the supplier's location, industry, or business size, if the supplied product or service affects AQWSR's deliverables to aerospace, defense, or regulated industries.
- Suppliers must ensure that all sub-tier providers understand and comply with these requirements and that controls are maintained throughout the supply chain.

# 2. Supplier Approval and Qualification

- All suppliers must be evaluated and approved by AQ Wiring Systems Rockford Ltd before receiving any
  purchase orders. This includes submission of a Supplier Questionnaire, quality system certifications (e.g., ISO
  9001, AS9100), and capability assessments.
- The supplier's approval status may be conditional or full, and is contingent upon risk level, product type, and criticality.
- AQWSR reserves the right to audit or inspect the supplier's facilities and processes as part of the qualification process.
- Suppliers must maintain their certification status and notify AQWSR immediately of any suspensions, revocations, or scope changes in their quality system certifications.
- Re-evaluation of approved suppliers will occur periodically, based on performance, audit results, and criticality of the products or services supplied.

#### 3. Flow Down of Requirements

- Suppliers are responsible for communicating AQWSR's requirements, including technical drawings, specifications, and regulatory mandates, to their sub-tier suppliers.
- Sub-tier suppliers must be held to the same standards outlined in this manual, and compliance must be verified by the primary supplier.
- Suppliers must ensure the traceability of flowed-down requirements, maintaining records of what was provided, when, and to whom.
- When AQWSR's customers impose specific requirements, these must also be flowed down to all relevant sub-tiers without omission.

## 4. Product and Service Conformity

- Suppliers must ensure that all products and services provided meet applicable AQWSR purchase order terms, specifications, drawings, industry standards, and customer-specific requirements.
- Products shall be free of damage, defects, and contamination. Packaging must protect the product from damage during handling and transportation.
- All materials must be verified for correct revision status, part numbers, and condition prior to shipment.
- Where applicable, Certificates of Conformance (CoCs), inspection reports, and supporting documentation must accompany the shipment.
- For shelf-life controlled products, the remaining shelf life at time of delivery must be at least 80% of the total unless otherwise authorized in writing by AQWSR.



# **5. Special Processes Control**

- Special processes include operations such as heat treatment, welding, plating, painting, anodizing, chemical
  processing, and any other process where the resulting output cannot be fully verified by subsequent
  inspection.
- Suppliers performing special processes must validate these processes through documented procedures, operator qualifications, and process controls.
- When applicable, suppliers shall maintain NADCAP accreditation or obtain specific written approval from AQWSR before conducting such processes.
- Records of special process qualifications, training records, and process monitoring results must be retained and made available upon request.

## 6. Key Characteristics and Critical Items

- Key characteristics are defined as features that have a significant impact on fit, function, performance, or compliance with regulatory requirements.
- Suppliers must identify key characteristics using drawings, specifications, or AQWSR directives and apply
  appropriate control methods such as Statistical Process Control (SPC) or 100% inspection.
- All deviations from control limits or tolerance thresholds must be investigated and corrective actions implemented to prevent recurrence.
- Documentation of inspection results, non-conformance analysis, and preventive measures must be maintained.

## 7. Change Management

- Suppliers shall not change any design, manufacturing process, test methods, materials, facility location, or suppliers without prior written approval from AQWSR.
- Engineering Change Requests (ECRs) must be submitted to AQWSR at least 30 calendar days before the intended implementation date.
- All proposed changes must include a complete description, reason for change, affected part numbers, proposed implementation date, and any supporting risk analysis or validation evidence.
- AQWSR will evaluate the change for potential impact and may require First Article Inspection (FAI),
   additional testing, or requalification before approving the change.
- No affected product shall be shipped until AQWSR has issued formal written approval for the change.

### 8. First Article Inspection (FAI)

- First Article Inspection (FAI) ensures the first production run of a new or revised part meets all design and manufacturing specifications.
- Suppliers must conduct FAI in accordance with AS9102 and submit the full FAIR package to AQWSR for approval prior to delivering production parts.
- FAI is required for new parts, changes to design, changes to manufacturing processes, and relocation of production facilities.
- The FAIR must include all dimensional results, material certifications, special process approvals, and test data as applicable.



# 9. Non-Conformance Management and Corrective Action

- Non-conforming products must be segregated, clearly labeled, and quarantined to prevent unintended shipment.
- If a supplier identifies or is notified of a non-conformance, they must initiate containment within one business day and provide a full 8D root cause analysis and corrective action plan within 10 business days.
- All concessions or requests to ship non-conforming material must be documented and approved in writing by AQWSR before shipment.
- AQWSR reserves the right to reject any non-conforming material, issue chargebacks for rework, and suspend supplier approval status for repeated or critical issues.
- Supplier Action Required: Suppliers must respond using 8D format with strict adherence to the stated response times.

Non-conforming products may not be shipped without written concession.

Suppliers must follow the 8D process with specified timelines for containment and corrective actions.

| Event Occurs   |                |
|--|----------------|
| Problem notification and submission of component to supplier by Company    | Starting Point |
| Formal acknowledgement by supplier that reported problem has been received | 1 Day          |
| Receipt of samples at supplier factory                                     | 2 Days         |
| Initial supplier problem verification complete and communicated            | 3 Days         |
| Interim containment plan communicated to Company                           | 5 Days         |
| Supplier failure or problem analysis completed, and results communicated   | 9 Days         |
| Supplier corrective action plan communicated                               | 10 Days        |
| Supplier corrective and preventive action implemented and verified         | As per plan    |

## 10. Audits and Right of Access

- AQ Wiring Systems Rockford Ltd, its customers, and regulatory authorities reserve the right to perform audits of the supplier's quality system, processes, and facilities with reasonable notice.
- Audit types include system audits, process audits, and product audits, and may be triggered by routine scheduling, poor performance indicators, or customer/regulatory requests.
- Suppliers must also enable AQWSR to access their sub-tier suppliers if deemed necessary for verification or investigation purposes.
- Audit findings will be discussed during the closing meeting and a formal report will be issued. Suppliers are expected to respond with corrective actions within specified timelines.
- All audit results are confidential unless otherwise agreed upon with the supplier.

# 11. Traceability and Counterfeit Prevention

- Suppliers must ensure complete traceability of components, materials, and sub-assemblies from origin to delivery. This includes linking back to the Original Equipment Manufacturer (OEM) or authorized distributor.
- Labels and documentation must clearly identify batch/lot numbers, date codes, part numbers, and trace codes.
- Traceability must be maintained even if production lots are split, re-batched, or repackaged, and sub-lots must be distinctly traceable and not mixed.
- Counterfeit part prevention requires documented procedures to verify the authenticity of all products supplied. Suppliers must only source materials from approved and reputable sources.



• Suspected counterfeit products must be reported to AQWSR immediately and isolated to prevent unintended use or shipment.

| Category/Type           | Traceability Requirements   |
|-------------------------|---|
| Raw Material            | Lot traceable to manufacturer's part no, lot no, date code with Certified Mill  |
|                         | Test Report or Material Test Report.  |
| Manufactured Parts      | Traceability shall be maintained for all product throughout production from     |
|                         | raw material to finished product (including product quantities, split orders,   |
|                         | nonconforming product etc.).  |
|                         | Raw material used shall be Lot traceable to manufacturers part no, lot no, date |
|                         | code with Certified Mill Test Report or Material Test Report.                   |
|                         | Requirements specified for COTS; MOTS & Mechanical/Electrical parts also        |
|                         | apply where applicable.   |
| Commercial of the Shelf | Traceability through supply chain to Original Equipment Manufacturer (OEM).     |
| (COTS)                  |   |
| Modified of the Shelf   | Traceability shall be maintained for all product modifications throughout       |
| (MOTS)                  | production from material to finished product (including product quantities,     |
|                         | split orders, nonconforming product etc.).                                      |
|                         | The originating COTS item(s) shall have Traceability through supply chain to    |
|                         | Original Equipment Manufacturer (OEM).  |
|                         | Requirements specified for Raw Material & Mechanical/Electrical parts also      |
|                         | apply where applicable for item modifications.                                  |
| Mechanical / Electrical | Lot traceable to Original Equipment Manufacturers part no, lot no, date code.   |
| Parts                   |   |

#### 12. Documentation and Record Retention

- Suppliers must retain all quality-related records, including inspection reports, material certifications, process validation data, and calibration records for a minimum of 13 years unless otherwise specified on the purchase order.
- Upon request, suppliers must provide AQWSR or its customers with access to relevant records within two working days.
- Disposition of records after the retention period must be managed securely to maintain confidentiality.

#### 13. Competence, Training, and Awareness

- Suppliers must ensure that all employees performing work affecting quality are competent based on education, training, skills, and experience.
- Training programs must be documented, and records must be maintained to demonstrate employee qualifications and ongoing competence.
- Personnel must be made aware of their contributions to product or service conformity, product safety, and the importance of ethical behavior.
- Suppliers must periodically evaluate training effectiveness and ensure updates for changes in processes, products, or customer requirements.
- When required, evidence of training and qualification must be provided to AQWSR upon request.
- Personnel must be trained and aware of their impact on quality, safety, and ethics.
- Training records must be maintained.



## 14. Performance Monitoring

- Supplier performance is monitored continuously through metrics such as product quality (PPM), on-time delivery, responsiveness to non-conformance issues, and audit findings.
- Suppliers may receive performance scorecards on a quarterly or annual basis, and trends will be used to determine risk levels and continued approval.
- Failure to meet performance thresholds may result in additional scrutiny, formal corrective actions, and potential removal from the approved supplier list.
- Suppliers are encouraged to monitor their own performance against AQWSR's expectations and initiate self-directed improvements.
- Supplier performance is continuously evaluated based on quality (PPM), on-time delivery, responsiveness to corrective actions, and audit outcomes.
- Key metrics include delivery adherence, defect rates, concession frequency, and responsiveness to investigations or change requests.
- Suppliers may receive periodic scorecards or report cards and are expected to address any areas of concern proactively.
- Consistently poor performance may result in supplier probation, increased oversight, removal from the approved list, or financial liability for non-conformances.

#### 15. Ethical and Sustainable Business Practices

- Suppliers are expected to uphold the highest standards of ethical conduct and legal compliance in all business interactions.
- Unethical practices such as bribery, corruption, collusion, falsification of records, and intellectual property theft are strictly prohibited.
- Suppliers must comply with all applicable labor laws, including those related to child labor, working hours, discrimination, and safe working conditions.
- AQWSR encourages suppliers to adopt and promote environmental sustainability practices such as waste reduction, pollution control, and responsible material sourcing.
- Audits or evaluations of supplier ethical practices may be conducted, and non-compliance may lead to suspension or termination of business relationships.
- Suppliers must comply with all applicable laws and regulations relating to human rights, labor practices, environmental impact, and ethical sourcing.
- Child labor, forced labor, and unsafe working conditions are strictly prohibited.
- Suppliers are expected to provide fair compensation, working hours within legal limits, and a workplace free of harassment and discrimination.
- Environmental sustainability practices, such as minimizing waste, pollution control, and responsible material sourcing, are strongly encouraged.
- AQWSR reserves the right to audit supplier's ethical practices and may require evidence of compliance as a condition of continued approval.

#### 16. Additional AS9100 Clause 8.4.3 Requirements

- Suppliers must ensure personnel are competent and, where applicable, qualified or certified to perform designated tasks.
- Statistical techniques must be applied where required for process or product validation, and documented acceptance criteria must be followed.
- Suppliers must provide test specimens for design validation, product verification, or auditing purposes when required by contract.



- Personnel must be made aware of how their work contributes to the conformity of products and services, as well as product safety.
- Suppliers must promote a culture of quality and compliance aligned with AS9100 Rev D expectations.
- Suppliers must demonstrate competence, maintain documentation, and provide test specimens when required.

## 17. Definitions and Terminology

FAIR: First Article Inspection Report per AS9102.

ECR: Engineering Change Request – a formal proposal for a product or process change.

Concession: Formal approval to accept a non-conforming product as-is or with rework.

PPM: Parts Per Million – a defect rate metric.

COTS: Commercial Off-The-Shelf parts.

MOTS: Modified Off-The-Shelf parts.

Zero Defects: A philosophy aimed at reducing defects to zero.

Counterfeit: Unauthorized copies, substitutions, or modified components misrepresented as genuine.