

QUALITY REQUIREMENTS

This Quality Assurance Specification establishes the specific requirements that apply when one or more of the following Quality Requirements (QR's) are specified by code on **INTERSTATE PLASTICS**Purchase Orders. These requirements are in addition to those set forth in any other contractual document. The provisions indicated herein are an integral part of the Purchase Order.

Compliance with these requirements does not reduce Seller responsibility for furnishing materials and services, which fully comply with all applicable Drawing(s) and Specification(s), nor does it guarantee acceptance of materials or services by **INTERSTATE PLASTICS**. In the event that materials or services are found to be defective and cannot be demonstrated by the Seller to be in conformance with the Purchase Order, **INTERSTATE PLASTICS** has the right to reject them.

Suppliers will be notified of any changes to these Quality Requirements.

INDEX

- QR 1 Inspection System
- QR 2 Control Of Changes
- QR 3 Supplier Corrective Action
- QR 4 Right Of Entry
- QR 5 Shelf Life And Temperature Sensitive Materials
- QR 6 Packaging
- QR 7 Material Safety Data Sheets
- QR 8 Identification
- QR 9 Calibration System Requirements
- QR 10 Certificate Of Conformance
- QR 11 Material Test Reports
- QR 12 Certified Test Data
- QR 13 Approved Processor Requirements
- QR 14 Report Of Discrepancy
- QR 14.1 Report Of Escape
- QR 14.2 Notification of Non-Compliance
- QR 15 Traceability/Lot And Batch Control
- QR 16 Supplier Records
- QR 17 Key Characteristics
- QR 18 Supplier Flow-down to Sub-supplier
- QR 19 AAM Requirements



1. INSPECTION SYSTEM REQUIREMENTS:

The Seller shall provide and maintain an inspection system in conformance with:

- NADCAP
- ISO 9001, latest revision
- AS9100, latest revision
- or other Quality System approved by Interstate Plastics.

2. CONTROL OF CHANGES:

Seller agrees not to make any change in materials or design details or other product which would affect the part or any component part thereof without prior written Buyer approval. The Seller will identify, on the Certificate of Conformance and/or packing sheet, the as built revision level of the end item product being delivered.

3. SUPPLIER CORRECTIVE ACTION:

Seller shall, on request, on forms designated by Buyer, provide statements of corrective action on failures of seller's hardware or quality system. Corrective action statements, at Buyer's option may require approval signature by Buyer and Government Quality representative. All rejected articles resubmitted by seller to Buyer shall bear adequate identification including reference to Buyer's rejection document.

4. RIGHT OF ENTRY:

(1) The Buyer, their customer and regulatory authorities shall be granted the right of acces to all seller's facilities involved in the order and all applicable quality records.

(2) The right to verify at the seller's premises that subcontracted product conforms to specified requirements. Such verification shall not be used by the seller as evidence of effective control of quality by the supplier.

5. SHELF LIFE AND TEMPERATURE SENSITIVE MATERIALS:

The Seller shall identify all materials and articles which have definite characteristics of quality degradation with age or environment. The Seller shall affix this information directly on the material container or article. This identification shall indicate the date useful life was initiated and the date or cycle at which the useful life will be expended. When environment is a factor in determining useful life, the identification shall include the storage condition (i.e., temperature, humidity, etc.) required to achieve the stated life. A minimum of 75% of the applicable material/article shell life shall remain upon receipt of the material by Buyer or the material is subject to rejection and returned to the seller.

6. PACKAGING:

Unless otherwise specified by the item drawing, specification or purchase order, the seller is responsible for assuring that all items are delivered without damage or deterioration and are efficiently and economically packed for the method of transportation and type of handling involved. Unit and intermediate packaging will be employed as necessary to prevent damage or deterioration.

7. SAFETY DATA SHEETS:

Materials(s) noted on this purchase order must be supplied in accordance with OSHA's hazard communication standard 29CFR1910-1200, OSHA instruction CPL2-2.38 dated May 10, 1998 and Washington State codes 296-62-05413. All first time orders MUST be supplied with "Safety Data Sheets". Materials not received in compliance with aforementioned OSHA requirements and WashingtonState codes will be subject to immediate rejection and return at supplier's expense.

In addition, if Seller is aware of any additional precautions and/or handling techniques instituted with regard to other customers, seller is required to requested to submit those safeguards with SDS.

Seller is required to forward a Toxic Substances Control Act (TSCA) certification letter to the Buyer for the product(s) purchased on this purchase order with the statement that every chemical component of the product(s) is listed by the Toxic Substances Control Act Inventory (P.L. 94-94-969).

8. IDENTIFICATION:

Parts, assemblies and components shall be

identified as specified on the engineering drawing. When identification is not specified on the engineering drawing, the

QUALITY REQUIREMENTS

product shall be identified with the part number specified on the purchase order. When items are too small to easily identify, they may be bagged and tagged with the proper identification indicated on the bag or tag. Raw material procured to Federal, Military, Aerospace or other specification shall be marked and identified per the requirements of the identification specification which is referenced in the controlling (i.e. Federal, Military, Aerospace, etc.) specification.

9. CALIBRATION SYSTEM REQUIREMENTS:

Seller shall have a calibration system that assures compliance with NIST / ANSI / NCSL Z540-1-1994 "American National Standard for calibration". Any deviation or waiver to this requirement must be approved by buyer's Material and Quality Assurance Representatives.

10. CERTIFICATE OF CONFORMANCE:

Each shipment will be accompanied by a legible and reproducible copy of a Certificate of Conformance with the signature of responsible representative stating material, process, or article being shipped meets requirements of applicable drawings or specification cited in Purchase Order. Supplier will include date of manufacture on Certificate of Conformance. If an Outsource Procurement Specification is called out on the Purchase Order, include the revision level.

11. MATERIAL TEST REPORTS:

A legible and reproducible copy of material test reports will accompany each shipment. Test reports will be identified with specification number and heat and/or cure lot number. Chemical and physical test reports will include actual numerical values for each property tested in accordance with the applicable specification. When more than one specimen is required, test results of each is required on the report. Specification and revision will be shown on each test report furnished.

12. CERTIFIED TEST DATA:

A legible and reproducible copy of certified test data will accompany each shipment of material, parts, or assemblies. Test will be identified with specification number and heat and/or cure lot number. Chemical and physical test data will include actual numerical values for each property tested in accordance with the applicable specification. Specification and revision will be shown on each report furnished. Test data will include the following statement (or equivalent): "Test reports are on file and available upon request."

13. APPROVED PROCESS REQUIREMENTS:

A legible and reproducible copy of special process certifications (i.e. testing, heat treat, nondestructive testing, etc.) will accompany each shipment of material, parts, or assemblies. Special processes will be performed by accredited process facilities and, if required by contract, customer (process specification owner) approved sources.

14. <u>REPORT OF DISCREPANCY:</u>

Any departure from drawing specifications or other purchase order requirements must be documented by the Seller and submitted to the Buyer for consideration and disposition. A copy of this disposition document must accompany each affected shipment.

14.1 REPORT OF ESCAPE:

The Seller upon discovery of a shipment / delivery of nonconforming product must provide a notice of disclosure to the Buyer within 48 hours of discovery.

14.2NOTIFICATION OF NON-COMPLIANCE

When an out-of-tolerance condition is discovered on a measuring device being calibrated by an approved Calibration Lab used by Interstate Plastics, that facility will contact Interstate Plastics immediately.



15. TRACEABILITY/LOST AND BATCH CONTROL:

Seller must maintain lot and batch control of raw materials to purchased items. Seller must provide positive traceability of manufactured parts and assemblies to raw materials through the use of lot/batch, serial numbers or date of manufacture, as applicable, for all items in the shipment.

16. SUPPLIER RECORDS:

The Seller shall maintain records of product delivered to buyer for a period of ten (10) years.

17. KEY CHARACTERISTICS:

The Seller shall maintain applicable statistical control charts for key characteristics identified by the customer drawings.

A copy of the for key characteristics statistical control charts must accompany all items in the shipment to Buyer.

18. SUPPLIER FLOW DOWN TO SUB-SUPPLIER

The Seller shall flow down to sub-tier suppliers the applicable requirements in the purchasing documents. In addition, Sub-tier suppliers are expected to conduct their business in an ethicalmanner, and with due regard to personal and product safety, and with a commitment to provide only genuine parts and materials. (Ref. definition of Counterfeit part in AS9120D.) **19.ACCEPTANCE AUTHORITY MEDIA:**

Supplier's quality systemshall include adequate controls of Acceptance Authority Media (AAM). This shall include but not be limited to:

Application Errors (i.e. Omission, Typos, Legibility, etc.)

• Untimely Use (i.e. Documentation is not completed as planned, "Stamp/Sign as you go", etc.)

• Misrepresentation (i.e. Uncertified personnel, Falsification of documentation, Work not performed as planned, etc.)

 Training Deficiencies (i.e. Ethics, Culture awareness, Proper use of authority media, etc.)

QUALITY REQUIREMENTS