

Quality Specification

GE Additive



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1 SCOPE AND BACKGROUND

1.1 Scope

This specification provides the General Quality Requirements for all GE Additive suppliers. This specification applies to all purchased products used in GE Additive.

The purpose of this Supplier Quality Requirements specification is to establish a set of procedures, practices and expectations pertaining to the quality of items purchased by GE Additive and qualification of Suppliers to GE Additive.

1.2 Language

“Shall” is used whenever a requirement expresses a provision that is mandatory.

“Will” is used to describe a task that is performed by an individual or organization not governed by the document. “Will” is not to be used to express a mandatory provision. Use the term “shall”.

“Should” and “may” are used when it is necessary to express non-mandatory provisions. When a non-mandatory provision is recommended “should” is used, otherwise “may” is used.

“Must” is not to be used to express a mandatory provision. Use the term “shall”. “Must” may be used in explanation to identify an external requirement or regulation that is to be met.

“Is” and “are” are used for descriptive text. No mandatory or non-mandatory requirements are expressed using these terms.

Throughout this specification the term “Purchaser” is intended to mean GE Additive and/or businesses acquired by GE Additive. The term “Supplier” is intended to mean suppliers and/or planned suppliers that will provide raw materials, parts and/or assemblies to GE Additive for consumption, performance of services, or for supply of GE Additive customers applicable documents.

1.3 Definitions

Build-to-Print Part – A part that is manufactured according to a GE drawing and associated GE specifications called out on that drawing.

Build-to-Specification (Non-Build-to-Print) Part – A part that is manufactured to meet the requirements of a GE functional specification rather than a GE drawing.

Containment – Actions taken to minimize or eliminate the risk to GE Additive or its customers associated with a nonconformance for product already produced or in process of being produced.

Corrective Action – Actions taken remove the causes of an existing nonconformity or undesirable situation on the next product produced.

Critical for Safety (CTS) – Those Characteristics of a process critical to safety.

Critical to Process Characteristic (CTP) – Those Characteristics of a process that combine to define a Critical to Quality Characteristic; or are deemed essential for quality assurance purposes.

Critical to Quality Characteristic (CTQ) – Those Characteristics of an item which if nonconforming, may prevent or seriously affect the unit performance, reliability, producibility, or customer satisfaction of a product.

Failure Mode and Effect Analysis (FMEA) - A systematic, proactive method for evaluating a process or system to identify where and how it might fail and to assess the relative impact of different failures.

Frozen Process – A manufacturing method, process, procedure or control that was approved by the GE Additive Qualification Team and may not be changed by the Supplier without prior approval of the GE Additive Qualification Team. The Qualification Team defines process steps that can be changed without approval as part of the Manufacturing Process Plan review and approval.

Manufacturing Process Plan (MPP) – A detailed, step-by-step sequence of operations and requirements by which products are manufactured.

Nonconformance – A product that does not comply with the Purchaser or Supplier specification or was produced outside of approved process, work instructions or procedures.

Non-Destructive Testing (NDT) – Analysis techniques used to evaluate properties of material, component or system without causing damage. Typical methods would include ultrasonic, magnetic-particle, liquid penetrant, radiography, eddy-current testing, etc.

Off-the-Shelf Products – Products are packaged solutions listed in a catalog which are then adapted to satisfy the needs of the purchasing organization.

Preventive Action – Action taken to eliminate the cause(s) of a potential nonconformance or undesirable potential situation to prevent occurrence of same or similar situations in the future.

Product Quality Plan (PQP) – A detailed, step-by-step list of operations and requirements in which a supplier identifies a process of how, what, why, when and who will perform tests or inspections and the applicable acceptance criteria. This may also be referred to as an Inspection and Test Plan (I.T.P.).

Product Safety Risk Assessment – Safety risk assessment for any supplier designed product in accordance with the principles defined by ISO 12100. Residual risk information should be provided to the GE Additive Qualification Team.

Qualification Package – All required documentation for a qualification. This may also be referred as Qualification Book.

Repair – A type of correction performed to a nonconformance that reduces but does not completely eliminate the nonconformance(s) such that the product is determined to be usable for its intended purpose.

Request for Design Change – A document submitted by the Supplier to request GE Additive Engineering's approval prior to implementing a change in design for the Supplier or its sub-tier supplier.

Rework – A type of correction performed to a nonconformance that completely eliminates the nonconformance(s) such that the product is determined to be conforming to specification or requirement in all respects.

Scrap – A disposition for nonconforming product that renders the material not useable for its intended purpose and/or that cannot be economically reworked or repaired in an acceptable manner.

Sourcing Representative – GE Additive representative who is authorized to negotiate price, delivery, terms and conditions, and place the Purchase Order for qualification and production. The Sourcing Representative owns communication with the Supplier for all commercial and fulfillment matters.

Supplier Deviation Request (SDR) – A request initiated by the Supplier to deviate from purchase order technical requirements (drawings, specifications, engineering instructions, etc.) or the approved qualification package.

Supplier Quality Engineer (SQE) – GE Additive representative who communicates the qualification requirements and is the key interface with the Supplier relative to qualifications, process improvements, nonconforming material dispositions, corrective actions, and surveillance auditing. SQE owns communication with the Supplier for all technical matters.

2 APPLICABLE DOCUMENTS

2.1 Issues of Documents

The following documents form a part of this specification to the extent specified herein. Unless otherwise indicated, the latest revision shall apply.

2.2 ISO Specifications

ISO 9001 – Quality Management System Requirements

ISO 12100 – Safety of machinery – General principles for design – Risk assessment and risk reduction

3 PROCESS

3.1 General Guidelines

It is the responsibility of the Supplier to define and implement a detailed quality system that ensures all products supplied to GE Additive are of the highest quality possible by conforming to GE Additive drawings and/or applicable specifications and meeting all the requirements set forth in this document.

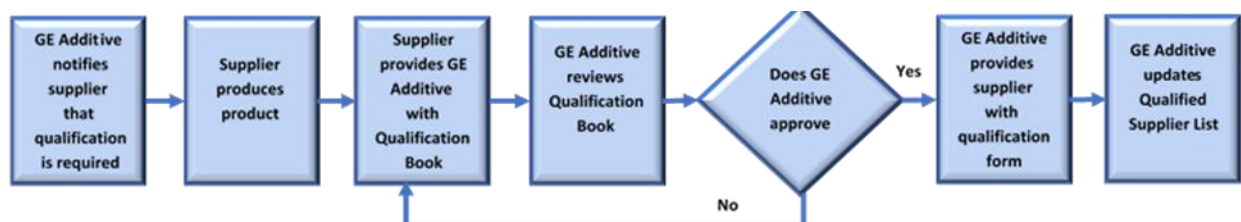
3.2 Communication

The GE Additive Purchase Order designates the Sourcing Representative who is the primary contact with the Supplier for commercial and fulfillment issues. The Supplier Quality Engineer (SQE) is the primary quality and technical contact and will be assigned by Supplier Quality management as appropriate.

3.3 Supplier Approval

In order to receive a GE Additive production purchase order, the Supplier shall be approved per the Know Your Supplier (KYS) process. Criteria for approval could include, but is not limited to, the following: properly executed confidentiality agreement (Non-Disclosure Agreement (NDA) or Proprietary Information Agreement (PIA)), acknowledgement of compliance with GE Additive Suppliers Integrity Guide, completion and passing of required technical capability assessment, EHS compliance/employment/security practices, financial viability, customer service aptitude, and strategic

Figure 1 – Qualification Process Flow Chart



value. The supplier approval process is performed prior to a qualification purchase order being issued to a supplier.

3.4 Supplier Qualification

After Supplier Approval is granted, GE Additive may require the Supplier to become qualified for each specific process, part or commodity family. If GE Additive requires the Supplier to perform a qualification, the SQE or Sourcing Representative will provide the Supplier with the Supplier Qualification Notification form in Appendix A. Through the qualification process, the Supplier demonstrates ability to provide high quality products on a repeatable basis in accordance with requirements and expectations of the GE Additive business purchasing the material. The qualification process applies to one product at one site and may pertain to certain pieces of equipment.

All Build-to Spec, Build-to-Print and Commercial off-the shelf product suppliers will be required to complete the qualification process, as described in this specification.

4 GENERAL REQUIREMENTS

4.1 Requirements

This section details the requirements that all Suppliers shall meet.

4.2 Quality System

The Supplier shall maintain a documented quality system to ensure control and conformance to the requirements of GE Additive's drawings and specifications. The quality management system shall meet the requirements of the current ISO 9001 (Quality management systems – Requirements) standard. Compliance to this requirement shall be demonstrated if requested by GE Additive by either of the following:

- Provision of a copy of a current certification(s) if requested, or
- Successful completion of a quality management systems audit to the current requirements of ISO 9001. GE Additive reserves the right to require this audit to be conducted by a third-party service designated by GE Additive or by a GE Additive representative. The Supplier will be responsible for all costs of the audit directly to the auditing party.

Any applicable industry standards (such as CE, UL, etc.) shall also be incorporated into the system. This system shall be made available to GE Additive for review upon request.

4.3 Record Retention

The Supplier shall have a written procedure for the documentation and retention of quality and product records for products supplied to GE Additive. The record retention period shall be a minimum of ten (10) years, from when the product was last shipped to GE Additive, unless otherwise specified by GE Additive or if a longer period is required by any applicable law or regulation. Records shall include, but are not limited to, product quality or inspection and test plans and results, material specifications, qualification documentation and certificates of conformance. Specific component record requirements may be specified in GE Additive purchase orders, contracts or specification. It is the responsibility of the Supplier to determine the appropriate storage means to meet the retention requirement and allow for timely retrieval of records.

4.4 Specification Transmittal to Suppliers

It is incumbent upon the Supplier to review with the Sourcing Representative and/or SQE the appropriate document retrieval/sharing methods that may be specific to their business. It is also the responsibility of the Supplier to review specification revisions with the Sourcing Representative and/or SQE on a continuous basis to ensure that the correct revisions are being worked to. When the Supplier receives a new purchase order, it is the Supplier's responsibility to verify they are working to the latest revision of the specification called out on the drawings and purchase order.

Unless otherwise notified by GE Additive, suppliers shall implement specification revisions on all existing and future purchase orders except where products have already entered the manufacturing process. Any exceptions to this policy shall be negotiated between the GE sourcing representative and Supplier.

If the Supplier does not have the latest revision of any relevant specification as described in the latest GE Additive purchase order, it is the Supplier's responsibility to request the latest revision of the specification.

4.5 Source Inspection and Test Witness

GE Additive and/or its customer may elect to inspect products, and/or witness subassemblies at the Supplier's facility during processing, testing, or at final inspection. All source inspection and test witness requirements will be identified and coordinated through the GE SQE, Quality Assurance, Quality representative or other designated representative. The Supplier shall provide all test samples.

4.5.1 Timing

The Supplier shall notify GE Additive in advance, when materials or products will be ready for inspection. The timing of this advance notification shall be a minimum of 2 weeks. GE Additive may decide to visit the Supplier facility.

4.5.2 Additional Requirements

Additional requirements for GE Additive and/or customer acceptance of product do not relieve the Supplier of its obligations to supply components that meet drawing and Purchase Order requirements.

4.6 Deviations

When a deviation to a requirement including a drawing, specification, MPP, packaging, or a material shortage is known or expected to exist, the Supplier shall submit a Supplier Deviation Request (SDR) as early in the process as possible to the SQE and Sourcing representative using form in Appendix C. If a deviation exists or could potentially exist, an SDR shall be submitted and approved prior to shipping the deviated products. The approved SDR applies to only the Purchase Orders listed on the SDR. SDRs shall be submitted by the Supplier for approval of alternate materials, processes, to correct drawing/documentation errors or omissions and other deviations to the PO requirements. SDRs shall be submitted by the primary Supplier (the Seller on the Purchase Order). Any deviations (e.g. drawing changes, material substitutions, etc.) related to a sub-tier supplier's scope that affect fit, form and/or function of the Supplier's product shall be submitted through the primary Supplier.

The SDR shall contain detailed description, containment, probable source and proposed remedial action (when business directed) information as part of the initial submittal. Failure to supply all of the information as required may result in the SDR being returned to the Supplier for completion of the required information. If this rejection impacts fulfillment requirements, charges may apply to the Supplier.

The Supplier shall not ship any deviating product before the SDR is approved by GE Additive. The Supplier shall send a copy of the approved SDR along with the product(s) at the time of shipment. GE has the right to request additional inspections and tests beyond applied drawing and specifications to prove the

deviated product's form, fit and function prior to SDR disposition. No repair shall be performed on a deviating product prior to disposition by GE.

In the case of nonconforming products or escaping defects, SDRs are "one-time" exceptions to GE Additive requirements. Unless the SDR involves a drawing change, GE Additive, expects the nonconformance(s) to be eliminated on subsequent deliveries.

Where appropriate, the Supplier shall provide a complete deviation description to include:

- Drawing/item number with zone of referenced area
- Inspection results
- Samples or photographs where applicable
- Number of defects for the lot(s) of material
- Specific Purchase Order numbers by product grouping
- Serial numbers of the components
- Estimated time to make correction(s)
- Cost related issues

For serialized parts, the serial number(s) shall be identified. For non-serialized parts, the specific Purchase Order(s) and date range shall be identified on the SDR.

For metal powder suppliers only: If a retest of any sample is determined to be nonconforming, the powder shall be rejected. In this case, the Supplier should not submit an SDR. GE Additive will not approve.

4.6.1 Containment

Containment is expected to be immediate when a nonconformance is discovered, with all products affected being contained. Containment actions apply to products, process and materials in which the nonconformance was detected as well as similar products or product families in which the nonconformance may occur. If the nonconformance is discovered during random audit, all WIP, inventory and shipped but not yet received products shall be evaluated.

Containment at the Supplier is expected to isolate (separate from normal production), insulate (inspect products to sort for defects at the Supplier, in transit for shipment and at the customer site) and aid in control of risk related to the nonconformance. An effective containment process documents the Supplier's efforts to verify control of its processes, (pre-production, production and post-production). The Supplier shall document and share all containment actions.

4.6.2 Probable Source

The Supplier shall report the source of the problem considering the following, as applicable:

- Situations involving the same or similar materials, products, equipment, instruments or system abnormalities and inconsistencies in the process that are also supplied to GE Additive
- Environmental conditions (e.g., temperature, humidity, light)
- Trends associated with equipment performance or specifications

- Cause code and deviation category

4.6.3 Proposed Remedial Action

Where applicable, Suppliers to GE Additive shall provide a rework or repair concept plan for all deviating products and services. Where rework or repair is not possible, substantiation shall be provided.

Rework or Repair Concept Plans shall include, as applicable:

- Identified risks that would adversely impact the product
- Planned completion date
- Estimated time (labor) required to complete correction

The Supplier shall have a positive identification plan, which ensures deviations and or corrected and or conforming materials are appropriately identified.

The Supplier shall document and show evidence to GE Additive that the remedial actions have been executed. GE Additive will evaluate whether the remediation execution eliminated the deviating condition or met the disposition requirements.

4.7 Corrective Action Procedures

When requested by the GE Additive SQE, the Supplier shall perform a formal root cause analysis (RCA) and identify containment, corrective, and preventive actions using the standard 8D RCA method & forms (or equivalent process). The Supplier shall provide an update on RCA to GE Additive within 2 weeks after it is requested and provide updates at least every 2 weeks until the RCA is completed. Corrective Action Requests that remain open longer than the specified period may result in disqualification of the Supplier.

Corrective action is intended to:

- Prevent the recurrence of the problem
- Avoid creation of further product or process issues

As a minimum, corrective action is necessary when:

- The production process is incapable or inconsistent
- A potential issue exists with machine, tooling, or fixtures

The Supplier shall provide and maintain objective evidence that the actions have been accomplished.

4.8 On-going Process Capability Checks

The Supplier shall, as a minimum, measure and record data for all critical to quality (CTQs) and critical to process (CTPs) characteristics identified on the drawings, specifications, through fixed process agreements between the supplier and GE Additive. The Supplier shall regularly analyze the CTQ and CTP data for process capability. When requested, the Supplier shall provide CTQ and CTP process capability reports to GE Additive within 5 business days. Under the direction of the SQE, the Supplier may be requested to execute improvement projects based on the process capability analysis.

5 QUALIFICATION REQUIREMENTS

5.1 Applicability

This section details the requirements that only Suppliers that are required to become Qualified must meet. As stated above, Suppliers that are required to complete qualification will be formally notified by Supplier Qualification Notification Form in Appendix A.

5.2 Qualification Book

The Supplier shall provide GE Additive with a qualification book that includes each of the below items that are applicable to the Supplier's product. If an item is not applicable to the Supplier's product, the supplier shall submit a Supplier Qualification Book Deviation Request (Appendix D) to the GE Additive SQE.

5.2.1 Sub-tier Suppliers

If a Supplier that is undergoing the qualification process (or is already qualified) chooses to outsource a critical process or purchase a critical component from another supplier, the Supplier shall perform a qualification and surveillance of all sub-tier suppliers in accordance with the GE Additive requirements listed in this specification or equivalent. GE Additive reserves the right to 1) review the Supplier's process for selection, qualification, and surveillance of sub-tier suppliers, 2) to approve, or disapprove, sub-tier supplier qualifications, 3) audit and monitor the sub-tier supplier's processes and facilities when deemed necessary. This requirement also applies if the Supplier is a sales representative or distributor that procures products that are supplied to GE Additive.

No.	Qualify Form Name	Comments
1	GE Purchase Orders	Provide copy of GE Purchase Orders for this project
2	GE Specifications/GE Drawings	Provide a list of all GE Specifications, and GE Drawings, including revision level. This includes, but not limited to: Ordering spec., Equipment/functional spec., General spec., Outline drawing copy, Piping and Instrumentation diagram copy, Electrical drawing copy.
3	Supplier Drawings/Specifications	Provide a copy of all Supplier generated drawings and specifications, including revision level. If the supplier chooses to use drawings or specifications published by industry standard organizations, these shall also be provided. In cases where Supplier IP is a concern, a list of drawings and specifications, including revision level is sufficient.
4	Supplier Manufacturing Process Plan (MPP)	Provide a copy of the Manufacturing Process Plan MPP is a detailed, step-by-step sequence of operations and requirements by which components or services are manufactured. A MPP shall, at a minimum contain the following information: (1) A list of all applicable GE drawings/specifications, ordering sheets, outline drawings, and special process documentation along with the latest revision letter/number. For build-to-specification parts, the Supplier shall provide a list of all Supplier/Industry standard drawings,

		<p>specifications and revisions.</p> <p>(2) Sequential listing of all operations and identification of Special Processes and associated procedures.</p> <p>(3) Critical to Quality steps.</p> <p>(4) Identification of all component parts and sources.</p> <p>(5) Identification of all critical sub-tier suppliers and their manufacturing locations. Critical sub-tier suppliers include but are not limited to Raw Material and any Special Process suppliers.</p> <p>(6) A sequence plan of all manufacturing and inspection steps with appropriate sign-off documentation. Supplier proprietary processes/documentation may be available for inspection/review by SQE and GE Engineering. The MPP shall refer to these documents by revision controlled document number in the MPP.</p> <p>(7) The manufacturing location.</p> <p>(8) Revision history.</p> <p>(9) A process flow diagram that shows the high-level process by which the product will be produced.</p> <p><i>For metal powder suppliers:</i> the MPP shall contain a contamination control plan. The contamination control plan shall include all major steps of powder production. Including:</p> <ul style="list-style-type: none">• Powder production• Material (or grade) changes• Powder handling• Air classification• Screening• Blending• Packaging• Ancillary Equipment <p><i>For metal powder suppliers:</i> the MPP shall identify the melting/atomizing practices utilized.</p>
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		<p><i>For metal powder suppliers:</i> The MPP shall identify the control process(es) for melting/atomizing (including atmosphere controls).</p> <p><i>For metal powder suppliers:</i> The MPP shall identify the process parameters for metal powder production.</p> <p>When the component is qualified, the MPP is considered part of the production Purchase Order requirements and may be frozen based on Qualification requirements.</p>
5	Product Quality Plan (PQP)/Inspection Test Plan (ITP)	<p>Provide a copy of the Supplier Product Quality Plan (PQP).</p> <p>The PQP shall, at a minimum, contain the following information:</p> <ul style="list-style-type: none"> • Clear identification of item, component, or system to which PQP is applicable (Drawing number(s) and product description(s)). • Listing of all technical documents that govern the inspection or test activity (i.e. Supplier documents, specifications, industry codes/standards) • Identification of the test or inspection criteria in an itemized listing. Each line item shall include: <ul style="list-style-type: none"> ○ What is to be inspected (to the characteristic level) ○ How it is to be inspected ○ What frequency it is to be inspected ○ When the inspection or test is to be performed (in manufacturing process) ○ Who is to perform the inspection (e.g., Operator, Inspector, etc.) ○ Acceptance criteria ○ Provision for sign off by the party performing the inspection • Sign-off documentation signifying completion of each inspection and test • Clear definition of customer involvement in the inspection and test activities (i.e. in-process inspections, customer witness and hold points, document reviews and and/or customer release inspections, etc.) • Identification and verification of CTQs and inspection methods.
6	Characteristic Accountability and Verification Forms (CAV)	<p>Provide a copy of the CAV report.</p> <p>The CAV form shall include, at a minimum, the following items:</p> <ol style="list-style-type: none"> (1) Identification of components (2) Characteristics and feature accountability (define critical parameters based on Design FMEA / e.g. dimensions, parallelism,) (3) Inspection and test results (4) Production Product Acceptance Criteria <p>Product acceptance criteria shall be established during the qualification process review of the CAV form. Once the level of inspection and product acceptance requirement has been determined and specified on the CAV form, it shall be applied to all production</p>

		<p>components hereafter to ensure controlled processes for maintaining drawing features and characteristics.</p> <p>A CAV form shall be completed and maintained by the Supplier.</p>
7	Measurement Method Validation	<p>Provide Measurement System Analysis (MSA) (ANOVA Gage R&R, Isoplot, etc...) for all critical measurements.</p> <p>Provide calibration certificates for all critical measurement tools.</p>
8	Bill of Materials (BOM)	<p>List to include Item #, description, drawing or specification number, revision, quantity, and serial numbers (in case of assembly of components).</p>
9	Component Conformance	<p>Include Certificate of Conformance for all critical components purchased from sub tiers.</p>
10	Design Calculations	<p>Where applicable, provide a copy of all design calculations for Components/Systems per Domestic and International Codes or regulation standards. This includes, but not limited to: Torque values, Material selection, Heat resistance, Lifetime, Dimensional fitting in assembly group.</p>
11	Product Safety Risk Assessment	<p>Safety risk assessment for any Supplier designed product in accordance with the principles defined by ISO 12100.</p>
12	Code Compliance	<p>Provide a copy of certificate for all Domestic and International Code Compliances that this product meets. Examples:</p> <ul style="list-style-type: none"> • IEC (International Electrotechnical Commission) • CE • ATEX • NEC (National Electrical Code) • UL
13	Material Test Reports	<p>Provide copies of Material Test Reports for all material used on this qualification/approval to include, but not limited to the following: Piping, Structural Steel, Bolting materials (Bolts, nuts, washers), Tubing, Raw Materials, Welding Consumables. Include all test reports for evaluation specifically called out as requirements or otherwise required for evaluating conformance to the drawings or specifications.</p>
14	Welding Procedures	<p>Only applicable if Supplier performs welding on GE Additive products.</p> <p>Provide a copy of the Welding/ Brazing Procedure, Specification, inspection, and all welder qualification records used on the Project.</p>
15	Surface Preparation and Painting	<p>Only applicable if Supplier performs painting on GE Additive products.</p>

		Include all metal preparation for, coating/paint procedures along with QA coating/paint data, signoffs, and coating/paint specifications
16	Sub-tier Supplier Qualification Documentation	All outsourced processes and purchased products that are incorporated into a product supplied to GE Additive require a qualification by first tier Supplier. Please provide MPP, PQP, MSA/gage R&R, Process FMEA (P-FMEA), audit report, etc. for each sub-tier supplier.
17	Functional Testing	Provide a copy of all Mechanical, Electrical, and Functional Tests performed. This should include testing procedures, documented data of all testing performed and test results. Example for one <u>complete</u> assembly (including serial number). SQE may require additional assessments over a lot of 10 or 20 complete assemblies.
18	Cleanliness	Provide a copy of the cleanliness procedure/work instruction used to verify cleanliness of product. <i>For metal powder suppliers:</i> the cleanliness procedures/work instructions shall include provisions to minimize opportunities for moisture pick-up and condensation.
19	Preservation and Packaging	Provide packaging proposal that includes how to preserve, crate, mark, and label the product (as applicable). Each package shall be labeled with the following information at a minimum (as approved by purchaser): <ul style="list-style-type: none"> • Supplier code • Part number with revision number • Box quantity • Box number • Manufacturing date • PO Number • Manufacturer • Serial Number • Lot Number • Safe Handling Instructions <i>For metal powder suppliers:</i> the packaging/preserving procedures/work instructions shall include provisions to minimize opportunities for moisture pick-up and condensation.
20	Repair/Rework	Provide any standard rework procedures that are applicable to the product(s) being supplied
21 ^{(M)(P)}	List of nonconformities	Provide a copy of list of production nonconformities (at Supplier or sub tier facility) related to this qualification/approval

22	Supplier Deviation Record List	Provide a copy or list of all SDRs used in this qualification/approval (Supplier and sub tier SDRs)
23	Photographs of the product or assembly	Provide photos of the completed product
24	Packing list	Provide a copy of the Packing List including, but not limited to, transport supports and sensors for transport monitoring.
25	GE-Certificate of Conformance	Provide a copy of Certificate of Conformance of the product. Evidence that the product was built according to current domestic and international standards.
26	Final Inspection Checklist	Provide a copy of the Final Inspection Checklist. This is the inspection checklist that Supplier will use every time before shipping a product or assembly to GE Additive. The expectation is that this is a live document that gets updated with additional checks based on lessons learned.
27	P-FMEA (Process Risk Assessment)	<p>Provide a copy of the P-FMEA.</p> <p>The P-FMEA relies on the results of the Design FMEA. The P-FMEA focuses on potential weaknesses in the production or performance process.</p> <p>When required by the qualification team, the Supplier shall perform a risk assessment of its manufacturing and quality assurance processes to evaluate the effectiveness of these processes to consistently produce the component or provide the qualified service. FMEA is one example of an accepted process risk assessment format</p>
28	Preventive Maintenance	Provide a copy of preventive maintenance plan for all equipment needed to manufacture product and that requires maintenance. This includes the calibration plan.
29	Other Documentation as requested	Any other documentation requested by the SQE

5.3 Qualification Approval Form

Upon successful completion of the qualification program and receipt of the Supplier Qualification Approval Form (Appendix B), or equivalent, the Supplier is released to fulfill subsequent Purchase Orders received from GE Additive. This qualification form indicates that, at the time of qualification and based on data provided by the Supplier, the manufacturing process used to produce the component(s) or perform a process was capable of complying with GE Additive drawing and specification requirements. No shipments are allowed before issuance of this form. Qualification approval does not relieve the Supplier of the full responsibility, on subsequent orders, to assure the manufacturing processes remain in control and the product or process supplied meets all drawing and specification requirements, unless formal, written approval for a deviation is obtained from GE Additive via a Supplier Deviation Request (SDR).

During the qualification, GE Additive and Supplier will define which steps of Manufacturing Process Plan (MPP) cannot be changed without approval of GE Additive. If changes are needed, a 'Supplier Deviation Form' shall be submitted by the Supplier before changes are implemented and wait for approval by GE Additive before implementing changes.

6 ADDITIONAL NOTES

6.1 Proprietary Nature of Information

As per the applicable confidentiality agreement, all information labelled as supplier proprietary and provided by the Supplier is held in confidence between GE and the respective Supplier and all GE information shall be held in confidence by the Supplier.

6.2 Supplier Integrity Guide

Link to the GE Suppliers Integrity Guide referenced in the Supplier Approval section above:
<http://www.gesupplier.com/html/SuppliersIntegrityGuide.htm>

APPENDIX A – SUPPLIER QUALIFICATION NOTIFICATION FORM

Supplier Qualification Notification Form

GE Additive requires the Supplier organization to complete formal supplier qualification in accordance with the requirements listed in P01AD502 for the below product(s). Please contact your Sourcing representative or Supplier Quality Engineering representative to acknowledge that you will comply with this request. Please respond within two weeks of receiving this form.

Notification

Date	Printed Name	Title	Signature and Date

Supplier Information

Supplier Name	
Supplier Site/Location	
Global Supplier Listing Number	

Product(s) Being Supplied

Raw Material/Part/Assembly/Machine Name	
Raw Material/Part/Assembly/Machine Number	
PO Number	

Supplier Acknowledgement

Function	Required	Printed Name	Title	Signature and Date
Sales	<input type="checkbox"/>			
Engineering	<input type="checkbox"/>			
Quality	<input type="checkbox"/>			

APPENDIX B – SUPPLIER QUALIFICATION FORM

Supplier Qualification Form

Supplier Information

Supplier Name	
Supplier Site/Location	
Global Supplier Listing Number	
Supplier Contact	
Supplier Contact Phone	
Supplier Contact Email	

Product(s) Being Supplied

Raw Material/Part/Assembly/Machine Name	
Raw Material/Part/Assembly/Machine Number	
PO Number	

Additional Information

Qualification Notes	
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Approvers

Function	Required	Printed Name	Signature and Date
Sourcing	<input type="checkbox"/>		
Engineering	<input type="checkbox"/>		
Materials Engineering	<input type="checkbox"/>		
Supplier Quality	<input type="checkbox"/>		

APPENDIX C – SUPPLIER DEVIATION REQUEST

Supplier Deviation Request

Supplier Information

Supplier Name	
Supplier Site/Location	
Global Supplier Listing Number	

Product(s) Being Supplied

Raw Material/Part/Assembly/Machine Name	
Raw Material/Part/Assembly/Machine Number	
PO Number	

Deviation Request

Specification Section	Original Requirement	Requested Requirement	Rationale

Additional Information

In the case of a nonconformance or escaping defect, provide the following (as applicable): detailed description, containment, probable source and proposed remedial action.

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Approvers

Function	Required	Printed Name	Signature and Date
Sourcing	<input type="checkbox"/>		
Engineering	<input type="checkbox"/>		
Materials Engineering	<input type="checkbox"/>		
Supplier Quality	<input type="checkbox"/>		

APPENDIX D – SUPPLIER QUALIFICATION BOOK DEVIATION REQUEST
SUPPLIER QUALIFICATION BOOK DEVIATION REQUEST

Supplier Information

Supplier Name	
Supplier Site/Location	
Global Supplier Listing Number	

Product(s) Being Supplied

Raw Material/Part/Assembly/Machine Name	
Raw Material/Part/Assembly/Machine Number	
PO Number	

Deviation Request

Qual Book Chapter	Rationale

Approvers

Function	Required	Printed Name	Signature and Date
Sourcing	<input type="checkbox"/>		
Engineering	<input type="checkbox"/>		
Materials Engineering	<input type="checkbox"/>		
Supplier Quality	<input type="checkbox"/>		

VERSION HISTORY

Version	Date (YYYY-MM-DD)	Issue Authority	Author
1	2020-02-12	AdEng-100233.A	A. Sprague
2	2020-07-07	AdEng-100253.A	A. Sprague