Identify in the matrix below where, in the quality system documentation, the information required in Section 2.0 of the ICC-ES Acceptance Criteria for Quality Documentation (AC10) can be found.

|  |  |
| --- | --- |
| Company Name: |       |
| Product/Material: |       |
| Evaluation Report, Listing or File No.: |       |
| Completed by:  |       | Date: |       |

| **AC10 SECTION** | **DOCUMENT IDENTIFICATION****AND PAGE NO.** | **DATE OF DOCUMENT** | **COMMENTS (IF NEEDED)** |
| --- | --- | --- | --- |
| **2.1.1****(Signature)** |       |       |       |
| **2.1.2****(Manufacturing location and contact info)** |       |       |       |
| **2.1.3****(Manual revisions)** |       |       |       |
| **2.1.4****(Product identification)** |       |       |       |
| **2.1.5****(Traceability)** |       |       |       |
| **2.1.6****(Work flow)** |       |       |       |
| **2.1.7****(Product changes)** |       |       |       |
| **2.1.8****(Organizational information)** |       |       |       |
| **2.1.9****(Packaging)** |       |       |       |
| **2.1.10****(Complaints procedure)** |       |       |       |
| **2.2.****(Incoming materials)** |       |       |       |
| **2.3****(In-process quality control)** |       |       |       |
| **2.4****(Final inspection)** |       |       |       |
| **2.5****(Nonconforming materials)** |       |       |       |
| **2.6.1****(Test equipment)** |      |       |      |
| **2.6.2****(Calibrations)** |       |       |       |
| **2.7.1****(QC forms)** |       |       |       |
| **2.7.2****(Document approval)** |       |       |       |
| **2.7.3****(Records retention)** |       |       |       |

|  |  |
| --- | --- |
| Signature: |       |
| Name of signer (type or print): |       |
| Name of third-party inspection agency (if applicable): |       |
| Date: |       |