Device Master Record (DMR)

Refer to *Procedure QP402: Device Master Record*

| Device name: | | | Type: | | Config: | |
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| Doc Code/No. | Document title | | Doc category | Type | Issued by | Date |
|  |  | | DS | Drawing | Dept/Function |  |
|  |  | | MP etc. | Specification | E.g. QA, Sales |  |
|  |  | |  | Procedure etc. |  |  |
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| **Approved by:**  <name>  <title> | | **Date:** | **Signature** | |  | |

Document Information

| Revision History | | | |
| --- | --- | --- | --- |
| Revision | Modified by | Change Control No. | Description of Change |
| 01 |  |  |  |
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Complete the above fields for each revision of this document. Ensure that there is sufficient description of changes so that the change history of this document can be followed. Additional columns can be added to include document/change tracking numbers generated by your company’s systems if required (eg. change control).

| Associated forms and procedures | |
| --- | --- |
| Doc. No. | Document Title |
| QP402 | Device Master Record |

List all controlled procedural documents referenced in this document (for example, policies, procedures, forms, lists, work/operator instructions

| Associated records | |
| --- | --- |
| Doc. No. | Document Title |
|  |  |

List all other referenced records in this document. For example, regulatory documents, in-house controlled documents (such as batch record forms, reports, methods, protocols), compliance standards etc.

DOCUMENT END