

Table 1.1–Documentation System Specifications

As specified in ORG 2.5.1, the Operator shall have a system for the management and control of documentation and/or data used directly in the conduct or support of operations. Such system shall comprise the elements specified below.

Note: Refer to the IRM for the definitions of Documentation, Electronic Documentation and Paper Documentation.

Elements		Documentation Types		
		Type 1	Type 2	Type 3
(i)	Identification of the version and effective date of relevant documents and/or data.	Recommended	Recommended	Required Note
(ii)	Identification of the title and, if applicable, sub-titles of relevant documents and/or data.	Recommended	Recommended	Required Note
(iii)	Distribution and/or dissemination that ensures all users are provided relevant documents and/or data on or before the effective date: (a) Throughout appropriate areas of the organization;	Required ^{Note}	Required ^{Note}	Required Note
	(b) To external service providers that conduct outsourced operational functions.			
(iv)	Definition of the specific media type(s) designated for presentation or display of the controlled version of relevant documents and/or data.	Required Note	Required Note	Required Note
(v)	Definition of documentation and/or data that is considered to be reproduced and/or obsolete.	Required Note	Required Note	Required Note
(vi)	Review and revision to maintain the currency of relevant documents and/or data.	Required Note	Required Note	Required Note
(vii)	Retention that ensures access to the content of relevant documents and/or data for a minimum period as defined by the Operator.	Required Note	Required Note	Required Note
(viii)	Provision for a scheduled backup by copying and archiving relevant documents and/or data, to include validation of the documents or data being backed up.	Required Note	Required Note	Required Note
(ix)	Identification and allocation of documentation access/user and modification rights.	Required Note	Required Note	Required Note
(x)	Dissemination and/or accessibility of documentation received from external sources such as regulatory authorities and original equipment manufacturers.	Required Note	Required Note	Required Note
(xi)	Identification of requirement for regulatory approval.	Required Note	Required Note	Required Note
Note: Required for conformity with ORG 2.5.1.				

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Standards and Recommended Practices

Table 1.2–Required Internal Audit Information

As specified in ORG 2.1.4, the Operator shall ensure the following information associated with the internal audit of individual ISARPs is recorded and retained:

- (i) The alpha-numeric identifier;
- (ii) Appropriate documentation reference(s) (from the Operator's documentation system);
- (iii) Auditor name(s);
- (iv) Audit date(s);
- (v) Auditor Actions accomplished by auditor(s) to provide evidence of implementation;
- (vi) If applicable, a description of non-conformance(s) and:
 - (a) The root cause(s) of non-conformance(s);
 - (b) The corrective action(s) implemented to address non-conformance(s).
- (vii) If applicable, a description of non-applicability (N/A);
- (viii) The current status of conformance (documented and implemented). GM

Note: The above-specified audit information may be retained in the Operator's electronic database as specified in ORG 2.1.4 and ORG 2.4.1, or in controlled procedural documents.

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