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Clause	Requirement	Comply	Auditor Notes / Evidence
	xt of the organization		
4.1 Unde	erstanding the organization and its context	1	
	The organization shall determine external and internal issues that are relevant to its purpose and its strategic direction and		
	that affect its ability to achieve the intended result(s) of its quality management system.		
	The organization shall monitor and review information about these external and internal issues.		
	NOTE 1 Issues can include positive and negative factors or conditions for consideration.		
	NOTE 1 Issues can include positive and negative factors of conditions for consideration. NOTE 2 Understanding the external context can be facilitated by considering issues arising from legal,		
	technological, competitive, market, cultural, social and economic environments, whether international, national, regional or		
	local.		
	NOTE 3 Understanding the internal context can be facilitated by considering issues related to values, culture,		
	knowledge and performance of the organization.		
4.2 Unde	erstanding the needs and expectations of interested parties		
	Due to their effect or potential effect on the organization's ability to consistently provide products and services that meet	1	
	customer and applicable statutory and regulatory requirements, the organization shall determine:		
	a) the interested parties that are relevant to the quality management system;		
	b) the requirements of these interested parties that are relevant to the quality management system.		
	The organization shall monitor and review information about these interested parties and their relevant requirements.		
4.3 Deter	rmining the scope of the quality management system		
	The organization shall determine the boundaries and applicability of the quality management system to establish its scope.		
	When determining this scope, the organization shall consider:		
	a) the external and internal issues referred to in <u>4.1;</u>		
	b) the requirements of relevant interested parties referred to in <u>4.2;</u>		
	c) the products and services of the organization.		
	The organization shall apply all the requirements of this International Standard if they are applicable within the determined		
	scope of its quality management system.		
	The scope of the organization's quality management system shall be available and be maintained as documented information. The scope shall state the types of products and services covered, and provide justification for any requirement		
	of this International Standard that the organization determines is not applicable to the scope of its quality management		
	system.		
	Conformity to this International Standard may only be claimed if the requirements determined as not being applicable do		
	not affect the organization's ability or responsibility to ensure the conformity of its products and services and the		
	enhancement of customer satisfaction.		
4.4 Quali	ity management system and its processes		
	<b>4.4.1</b> The organization shall establish, implement, maintain and continually improve a quality management		
	system, including the processes needed and their interactions, in accordance with the requirements of this International Standard.		
	The organization shall determine the processes needed for the quality management system and their application throughout the organization, and shall:		
	a) determine the inputs required and the outputs expected from these processes;		
	b) determine the sequence and interaction of these processes;		
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<ul> <li>c) determine and apply the criteria and methods (including monitoring, measurements and related performance indicators) needed to ensure the effective operation and control of these processes;</li> <li>d) determine the resources needed for these processes and ensure their availability;</li> <li>e) assign the responsibilities and authorities for these processes;</li> <li>f) address the risks and opportunities as determined in accordance with the requirements of <u>6.1;</u></li> <li>g) evaluate these processes and implement any changes needed to ensure that these processes achieve their intended results;</li> <li>h) improve the processes and the quality management system.</li> <li>4.4.2 To the extent necessary, the organization shall:</li> <li>a) maintain documented information to support the operation of its processes;</li> <li>b) retain documented information to have confidence that the processes are being carried out as planned.</li> </ul>	
5 Leadership	
5.1 Leadership and commitment	
5.1.1 General	
<ul> <li>Top management shall demonstrate leadership and commitment with respect to the quality management system by:</li> <li>a) taking accountability for the effectiveness of the quality management system;</li> <li>b) ensuring that the quality policy and quality objectives are established for the quality management system and are compatible with the context and strategic direction of the organization;</li> <li>c) ensuring the integration of the quality management system requirements into the organization's business processes;</li> <li>d) promoting the use of the process approach and risk-based thinking;</li> <li>e) ensuring that the resources needed for the quality management system are available;</li> <li>f) communicating the importance of effective quality management and of conforming to the quality management system requirements;</li> <li>g) ensuring that the quality management system achieves its intended results;</li> <li>h) engaging, directing and supporting persons to contribute to the effectiveness of the quality management system;</li> <li>i) promoting improvement;</li> <li>j) supporting other relevant management roles to demonstrate their leadership as it applies to their areas of responsibility.</li> <li><i>NOTE</i> Reference to "business" in this International Standard can be interpreted broadly to mean those activities that are core to the purposes of the organization's existence, whether the organization is public, private, for profit or not for profit.</li> </ul>	
5.1.2 Customer focus	
<ul> <li>Top management shall demonstrate leadership and commitment with respect to customer focus by ensuring that:</li> <li>a) customer and applicable statutory and regulatory requirements are determined, understood and consistently met;</li> <li>b) the risks and opportunities that can affect conformity of products and services and the ability to enhance customer satisfaction are determined and addressed;</li> <li>c) the focus on enhancing customer satisfaction is maintained.</li> </ul>	
5.2 Policy	· · · · · · · · · · · · · · · · · · ·
5.2.1 Developing the quality policy	
<ul> <li>Top management shall establish, implement and maintain a quality policy that:</li> <li>a) is appropriate to the purpose and context of the organization and supports its strategic direction;</li> <li>b) provides a framework for setting quality objectives;</li> </ul>	



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	c) includes a commitment to satisfy applicable requirements;	
5.2.2 Co	<ul> <li>d) includes a commitment to continual improvement of the quality management system.</li> <li>mmunicating the quality policy</li> </ul>	
5.2.2 60	The quality policy shall:	
	a) be available and be maintained as documented information;	
	b) be communicated, understood and applied within the organization;	
5.3 Orga	c) be available to relevant interested parties, as appropriate. nizational roles, responsibilities and authorities	
5.5 Orga		
	Top management shall ensure that the responsibilities and authorities for relevant roles are assigned, communicated and understood within the organization.	
	Top management shall assign the responsibility and authority for:	
	a) ensuring that the quality management system conforms to the requirements of this International Standard;	
	b) ensuring that the processes are delivering their intended outputs;	
	c) reporting on the performance of the quality management system and on opportunities for improvement (see	
	10.1), in particular to top management;	
	d) ensuring the promotion of customer focus throughout the organization;	
	<ul> <li>ensuring that the integrity of the quality management system is maintained when changes to the quality management</li> </ul>	
	system are planned and implemented.	
6 Planni		
	ons to address risks and opportunities	
U.I ACIN	<b>6.1.1</b> When planning for the quality management system, the organization shall consider the issues referred to in	
	4.1 and the requirements referred to in 4.2 and determine the risks and opportunities that need to be addressed to:	
	a) give assurance that the quality management system can achieve its intended result(s);	
	b) enhance desirable effects:	
	c) prevent, or reduce, undesired effects;	
	d) achieve improvement.	
	6.1.2 The organization shall plan:	
	a) actions to address these risks and opportunities;	
	b) how to:	
	<ol> <li>integrate and implement the actions into its quality management system processes (see <u>4.4</u>);</li> </ol>	
	<ol> <li>evaluate the effectiveness of these actions.</li> </ol>	
	Actions taken to address risks and opportunities shall be proportionate to the potential impact on the conformity of	
	products and services.	
	NOTE 1 Options to address risks can include avoiding risk, taking risk in order to pursue an opportunity, eliminating the	
	risk source, changing the likelihood or consequences, sharing the risk, or retaining risk by informed decision.eliminating	
	the risk source, changing the likelihood or consequences, sharing the risk, or retaining risk by informed decision.	
	NOTE 2 Opportunities can lead to the adoption of new practices, launching new products, opening new markets,	
	addressing new clients, building partnerships, using new technology and other desirable and viable possibilities to address	
	the organization's or its customers' needs.	
6.2 Qual	ity objectives and planning to achieve them	
UL GUU	<b>6.2.1</b> The organization shall establish quality objectives at relevant functions, levels and processes needed for the	
	quality management system.needed for the quality management system.	
	The quality objectives shall:	
	a) be consistent with the quality policy;	
	b) be measurable:	
I	b) be measurable,	1



<li>c) take into account a</li>	pplicable requirements;
d) be relevant to con	ormity of products and services and to enhancement of customer satisfaction;
e) be monitored;	
f) be communicated;	
g) be updated as ap	ropriate.
-,	maintain documented information on the quality objectives.
-	how to achieve its quality objectives, the organization shall determine:
	now to achieve its quality objectives, the organization shall determine.
,	
b) what resources wi	
c) who will be respon	
d) when it will be cor	
e) how the results wi	I be evaluated.
6.3 Planning of changes	
ů, s	determines the need for changes to the quality management system, the changes shall be carried
out in a planned mann	
The organization shall	consider:
a) the purpose of the	changes and their potential consequences;
b) the integrity of the	quality management system;
c) the availability of r	esources;
	allocation of responsibilities and authorities.
7 Support	
7.1 Resources	
7.1.1 General	
The organization sh	all determine and provide the resources needed for the establishment, implementation,
ũ	nual improvement of the quality management system. The organization shall consider:
a) the capabilities of	and constraints on, existing internal resources;
	bbtained from external providers.
7.1.2 People	
	determine and provide the persons necessary for the effective implementation of its quality
Ű	nd for the operation and control of its processes.
7.1.3 Infrastructure	
-	determine, provide and maintain the infrastructure necessary for the operation of its processes and
-	f products and services.
	ure can include:
a) buildings and asso	
	g hardware and software;
c) transportation resol	Irces;
d) information and cor	nmunication technology.
7.1.4 Environment for the ope	ration of processes
The organization shall	determine, provide and maintain the environment necessary for the operation of its processes and
to achieve conformity of	f products and services.
NOTE A suitable	environment can be a combination of human and physical factors, such as:
a) social (e.g. non-dis	criminatory, calm, non-confrontational);
<i>,</i>	stress-reducing, burnout prevention, emotionally protective);
b) psychological (e.g.	stress-reducing, burnout prevention, emotionally protective); erature, heat, humidity, light, airflow, hygiene, noise).
b) psychological (e.g. c) physical (e.g. temp	erature, heat, humidity, light, airflow, hygiene, noise).
b) psychological (e.g. c) physical (e.g. temp	erature, heat, humidity, light, airflow, hygiene, noise). r substantially depending on the products and services provided.

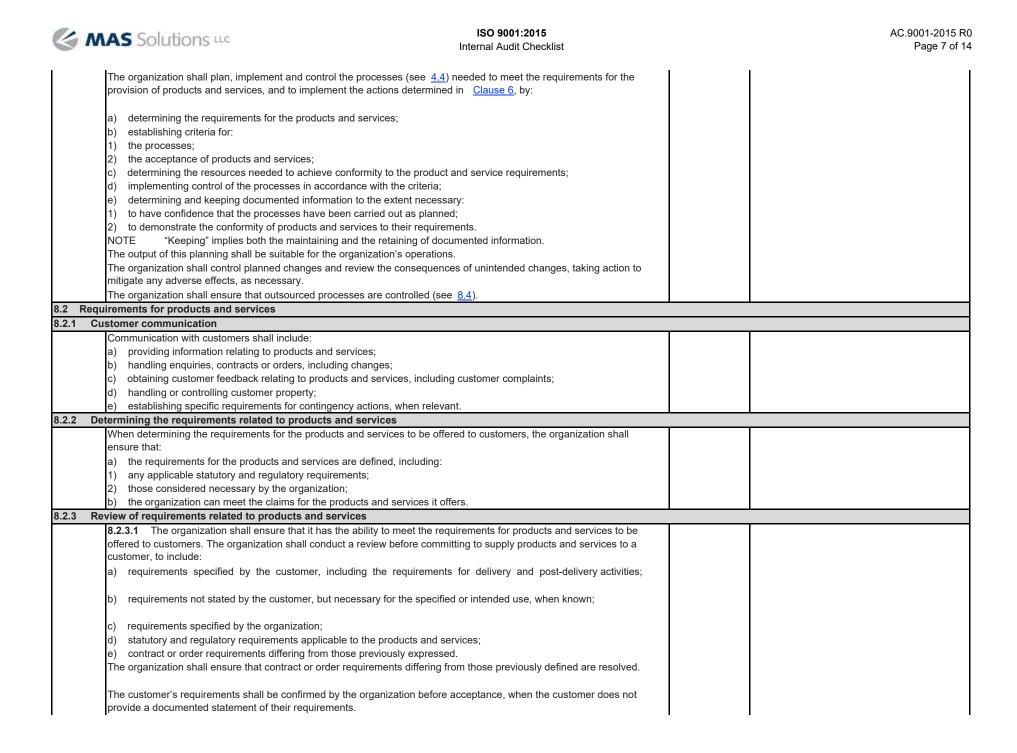


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7.1.5.1	General	
	The organization shall determine and provide the resources needed to ensure valid and reliable results when monitoring or	
	measuring is used to verify the conformity of products and services to requirements. The organization shall ensure that the	
	resources provided:	
	a) are suitable for the specific type of monitoring and measurement activities being undertaken;	
	<ul> <li>b) are maintained to ensure their continuing fitness for their purpose.</li> </ul>	
	The organization shall retain appropriate documented information as evidence of fitness for purpose of the monitoring and	
7.1.5.2	measurement resources. Measurement traceability	
1.1.3.2	When measurement traceability is a requirement, or is considered by the organization to be an essential part of providing	
	confidence in the validity of measurement results, measuring equipment shall be:	
	a) calibrated or verified, or both, at specified intervals, or prior to use, against measurement standards traceable to	
	international or national measurement standards; when no such standards exist, the basis used for calibration or	
	verification shall be retained as documented information;	
	b) identified in order to determine their status;	
	c) safeguarded from adjustments, damage or deterioration that would invalidate the calibration status and subsequent	
	measurement results.	
	The organization shall determine if the validity of previous measurement results has been adversely affected when	
	measuring equipment is found to be unfit for its intended purpose, and shall take appropriate action as necessary.	
	incusioning equipment is round to be unit for its interface purpose, and shall take appropriate action as necessary.	
7.1.6 O	rganizational knowledge	
	The organization shall determine the knowledge necessary for the operation of its processes and to achieve conformity of	
	products and services.	
	This knowledge shall be maintained and be made available to the extent necessary.	
	When addressing changing needs and trends, the organization shall consider its current knowledge and determine how to	
	acquire or access any necessary additional knowledge and required updates.	
	NOTE 1 Organizational knowledge is knowledge specific to the organization; it is gained by experience. It is information	
	that is used and shared to achieve the organization's objectives.	
	NOTE 2 Organizational knowledge can be based on:	
	a) internal sources (e.g. intellectual property; knowledge gained from experience; lessons learned from failures	
	and successful projects; capturing and sharing undocumented knowledge and experience; the results of improvements in	
	processes, products and services);	
	b) external sources (e.g. standards; academia; conferences; gathering knowledge from customers or external providers).	
7.2 Com	npetence	
	The organization shall:	
	a) determine the necessary competence of person(s) doing work under its control that affects the performance and	
	effectiveness of the quality management system;	
	b) ensure that these persons are competent on the basis of appropriate education, training, or experience;	
	c) where applicable, take actions to acquire the necessary competence, and evaluate the effectiveness of the actions	
	taken;	
	d) retain appropriate documented information as evidence of competence.	
	NOTE Applicable actions can include, for example, the provision of training to, the mentoring of, or the re-	
	assignment of currently employed persons; or the hiring or contracting of competent persons.	
7.3 Awa		
	The organization shall ensure that persons doing work under the organization's control are aware of:	
	a) the quality policy;	
	b) relevant quality objectives;	
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c) their contribution to the effectiveness of the quality management system, including the benefits of improved		
performance;		
d) the implications of not conforming with the quality management system requirements.		
7.4 Communication	-	•
The organization shall determine the internal and external communications relevant to the quality management system,		
including:		
a) on what it will communicate:		
b) when to communicate:		
c) with whom to communicate;		
d) how to communicate;		
e) who communicates.		
7.5 Documented information		
7.5.1 General		
The organization's quality management system shall include:		
a) documented information required by this International Standard;		
b) documented information determined by the organization as being necessary for the effectiveness of the quality		
management system.		
NOTE The extent of documented information for a quality management system can differ from one		
organization to another due to:		
<ul> <li>the size of organization and its type of activities, processes, products and services;</li> </ul>		
<ul> <li>the complexity of processes and their interactions;</li> </ul>		
— the competence of persons.		
7.5.2 Creating and updating		
When creating and updating documented information, the organization shall ensure appropriate:		
a) identification and description (e.g. a title, date, author, or reference number);		
b) format (e.g. language, software version, graphics) and media (e.g. paper, electronic);		
c) review and approval for suitability and adequacy.		
7.5.3 Control of documented information		
7.5.3.1 Documented information required by the quality management system and by this International Standard shall be		
controlled to ensure:		
a) it is available and suitable for use, where and when it is needed;		
b) it is adequately protected (e.g. from loss of confidentiality, improper use, or loss of integrity).		
7.5.3.2 For the control of documented information, the organization shall address the following activities, as		
applicable:		
a) distribution, access, retrieval and use;		
b) storage and preservation, including preservation of legibility;		
c) control of changes (e.g. version control);		
d) retention and disposition.		
Documented information of external origin determined by the organization to be necessary for the planning and operation		
of the quality management system shall be identified as appropriate, and be controlled.		
Documented information retained a sevidence of conformity shall be protected from unintended alterations.		
NOTE Access can imply a decision regarding the permission to view the documented information only, or the		
permission and authority to view and change the documented information.		
8 Operation		
8.1 Operational planning and control		





	NOTE In some situations, such as internet sales, a formal review is impractical for each order. Instead, the review		
	can cover relevant product information, such as catalogues or advertising material.		
	<b>8.2.3.2</b> The organization shall retain documented information, as applicable:		
	a) on the results of the review;		
	b) on any new requirements for the products and services.		
8.2.4	Changes to requirements for products and services		
	The organization shall ensure that relevant documented information is amended, and that relevant persons are made		
	aware of the changed requirements, when the requirements for products and services are changed.		
0.0 D			
	esign and development of products and services		
8.3.1	General		-
	The organization shall establish, implement and maintain a design and development process that is		
	appropriate to ensure the subsequent provision of products and services.		
8.3.2	Design and development planning		
0.0.1			
	In determining the stages and controls for design and development, the organization shall consider:		
	<ul> <li>a) the nature, duration and complexity of the design and development activities;</li> </ul>		
	<li>b) the required process stages, including applicable design and development reviews;</li>		
	c) the required design and development verification and validation activities;		
	d) the responsibilities and authorities involved in the design and development process;		
	,		
	e) the internal and external resource needs for the design and development of products and services;		
	<ul> <li>f) the need to control interfaces between persons involved in the design and development process;</li> </ul>		
	<ul> <li>g) the need for involvement of customers and users in the design and development process;</li> </ul>		
	h) the requirements for subsequent provision of products and services;		
	i) the level of control expected for the design and development process by customers and other relevant interested		
	parties;		
	j) the documented information needed to demonstrate that design and development requirements have been met.		
8.3.3	Design and development inputs		
	The organization shall determine the requirements essential for the specific types of products and		
	services to be designed and developed. The organization shall consider:		
	a) functional and performance requirements;		
	, , , , , , , , , , , , , , , , , , , ,		
	<li>b) information derived from previous similar design and development activities;</li>		
	c) statutory and regulatory requirements;		
	d) standards or codes of practice that the organization has committed to implement;		
	e) potential consequences of failure due to the nature of the products and services.		
	Inputs shall be adequate for design and development purposes, complete and unambiguous. Conflicting design and		
	development inputs shall be resolved.		
	The organization shall retain documented information on design and development inputs.		
8.3.4	Design and development controls		
	The organization shall apply controls to the design and development process to ensure that:		
1	a) the results to be achieved are defined:		
	b) reviews are conducted to evaluate the ability of the results of design and development to meet requirements;		
1	b) reviews are conducted to evaluate the ability of the results of design and development to meet requirements,		
1	c) verification activities are conducted to ensure that the design and development outputs meet the input requirements;		
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	d) validation activities are conducted to ensure that the resulting products and services meet the requirements for the		
	specified application or intended use;		
I	specified application of interfued use,	1	1



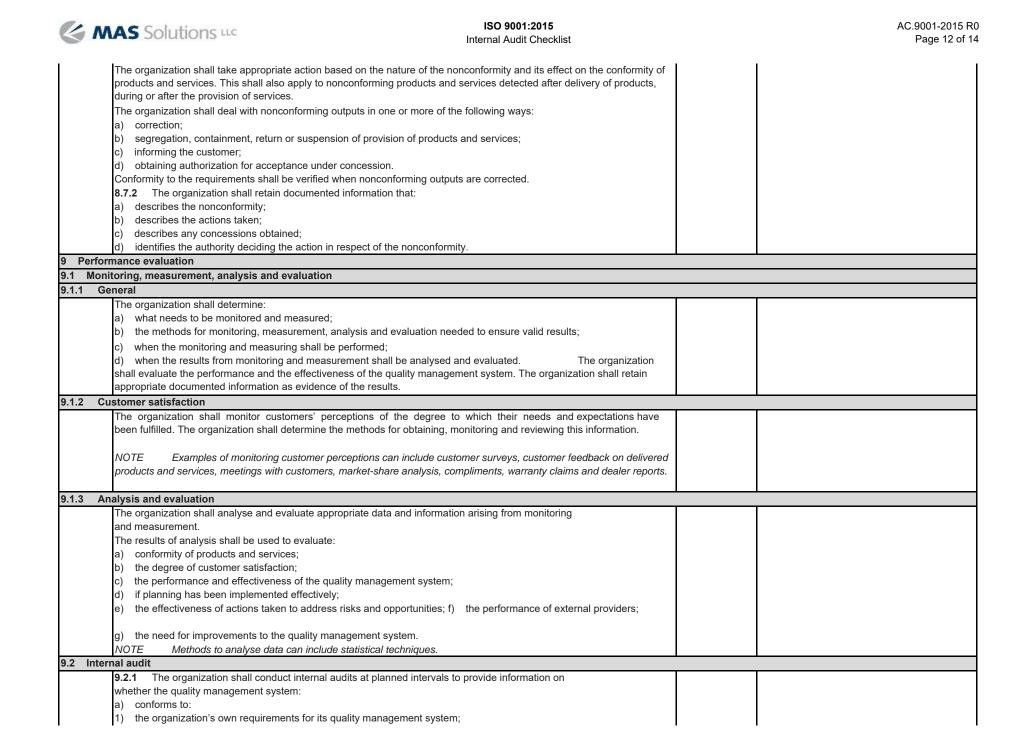
	e) any necessary actions are taken on problems determined during the reviews, or verification and validation activities;	
	f) documented information of these activities is retained.	
	NOTE Design and development reviews, verification and validation have distinct purposes. They can be conducted	
	separately or in any combination, as is suitable for the products and services of the organization.	
8.3.5	Design and development outputs	
	The organization shall ensure that design and development outputs:	
	a) meet the input requirements;	
	b) are adequate for the subsequent processes for the provision of products and services;	
	c) include or reference monitoring and measuring requirements, as appropriate, and acceptance criteria;	
	d) specify the characteristics of the products and services that are essential for their intended purpose and their safe and	
	proper provision.	
0.2.0	The organization shall retain documented information on design and development outputs.	
8.3.6	Design and development changes The organization shall identify, review and control changes made during, or subsequent to, the design and development of	
	The organization shall identify, review and control changes made during, or subsequent to, the design and development of products and services, to the extent necessary to ensure that there is no adverse impact on conformity to requirements.	
	products and services, to the extent necessary to ensure that there is no adverse impact on comornity to requirements.	
	The organization shall retain documented information on:	
	a) design and development changes;	
	b) the results of reviews;	
	c) the authorization of the changes;	
	d) the actions taken to prevent adverse impacts.	
8.4 C	ontrol of externally provided processes, products and services	
8.4.1	General	
	The organization shall ensure that externally provided processes, products and services conform to requirements.	
	The organization shall determine the controls to be applied to externally provided processes, products and services when:	
	a) products and services from external providers are intended for incorporation into the organization's own products and	
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		-	
	b) define both the controls that it intends to apply to an external provider and those it intends to apply to the resulting		
	output;		
	c) take into consideration:		
	1) the potential impact of the externally provided processes, products and services on the organization's ability to		
	consistently meet customer and applicable statutory and regulatory requirements;		
	consistentity meet customer and applicable statutory and regulatory requirements,		
	<ol> <li>the effectiveness of the controls applied by the external provider;</li> </ol>		
	d) determine the verification, or other activities, necessary to ensure that the externally provided processes, products		
	and services meet requirements.		
8.4.3	Information for external providers	-	
	The organization shall ensure the adequacy of requirements prior to their communication to the external provider.		
	The organization shall communicate to external providers its requirements for:		
	a) the processes, products and services to be provided;		
	b) the approval of:		
	1) products and services;		
	2) methods, processes and equipment;		
	3) the release of products and services;		
	c) competence, including any required qualification of persons;		
	d) the external providers' interactions with the organization;		
	e) control and monitoring of the external providers' performance to be applied by the organization;		
	f) verification or validation activities that the organization, or its customer, intends to perform at the external providers'		
	premises.		
0.5 D.			
	oduction and service provision		
8.5.1	Control of production and service provision	1	
	The organization shall implement production and service provision under controlled conditions.		
	The organization on an implement production and control provident and of contaitoned contaitions.		
	Controlled conditions shall include, as applicable:		
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1	The organization shall control the unique identification of the outputs when traceability is a requirement, and shall retain	1	1
	the documented information necessary to enable traceability.		
8.5.3	Property belonging to customers or external providers		
	The organization shall exercise care with property belonging to customers or external providers while it is under the		
	organization's control or being used by the organization.		
	The organization shall identify, verify, protect and safeguard customers' or external providers' property provided for use or		
	incorporation into the products and services.		
	When the property of a customer or external provider is lost, damaged or otherwise found to be unsuitable for use, the organization shall report this to the customer or external provider and retain documented information on what has		
	occurred.		
	NOTE A customer's or external provider's property can include material, components, tools and equipment,		
	premises, intellectual property and personal data.		
8.5.4	Preservation		
	The organization shall preserve the outputs during production and service provision, to the extent necessary to ensure		
	conformity to requirements.		
	NOTE Preservation can include identification, handling, contamination control, packaging, storage,		
	transmission or transportation, and protection.		
8.5.5	Post-delivery activities	-	
	The organization shall meet requirements for post-delivery activities associated with the products and services.		
	In determining the extent of post-delivery activities that are required, the organization shall consider:		
	a) statutory and regulatory requirements;		
	b) the potential undesired consequences associated with its products and services;		
	c) the nature, use and intended lifetime of its products and services;		
	d) customer requirements;		
	e) customer feedback.		
	NOTE Post-delivery activities can include actions under warranty provisions, contractual obligations such as		
950	maintenance services, and supplementary services such as recycling or final disposal.		
8.5.6	Control of changes The organization shall review and control changes for production or service provision, to the extent necessary to ensure		1
	continuing conformity with requirements.		
	The organization shall retain documented information describing the results of the review of changes, the person(s)		
	authorizing the change, and any necessary actions arising from the review.		
8.6 F	Release of products and services		
	The organization shall implement planned arrangements, at appropriate stages, to verify that the product and service requirements have been met.		
	The release of products and services to the customer shall not proceed until the planned arrangements have been		
	satisfactorily completed, unless otherwise approved by a relevant authority and, as applicable, by the customer.		
	The organization shall retain documented information on the release of products and services. The documented information shall include:		
	a) evidence of conformity with the acceptance criteria;		
	b) traceability to the person(s) authorizing the release.		
8.7 0	Control of nonconforming outputs	1	1
	<b>8.7.1</b> The organization shall ensure that outputs that do not conform to their requirements are identified and controlled to prevent their unintended use or delivery.		
I	to prevent their drimtended use of delivery.		I





2) the requirements of this International Standard;		
b) is effectively implemented and maintained.		
9.2.2 The organization shall:		
a) plan, establish, implement and maintain an audit programme(s) including the frequency, methods, responsibilities,		
planning requirements and reporting, which shall take into consideration the importance of the processes concerned,		
changes affecting the organization, and the results of previous audits;		
b) define the audit criteria and scope for each audit;		
<li>c) select auditors and conduct audits to ensure objectivity and the impartiality of the audit process;</li>		
<ul> <li>ensure that the results of the audits are reported to relevant management;</li> </ul>		
e) take appropriate correction and corrective actions without undue delay;		
f) retain documented information as evidence of the implementation of the audit programme and the audit results.		
NOTE See ISO 19011 for guidance.		
9.3 Management review		
9.3.1 General		
Top management shall review the organization's quality management system, at planned intervals, to ensure its		
continuing suitability, adequacy, effectiveness and alignment with the strategic direction of the organization.		
9.3.2 Management review inputs		
The management review shall be planned and carried out taking into consideration:		
a) the status of actions from previous management reviews;		
b) changes in external and internal issues that are relevant to the quality management system;		
c) information on the performance and effectiveness of the quality management system, including trends in:		
o/ montation on the performance and encourceded of the quarky management cyclem, monading action in		
1) customer satisfaction and feedback from relevant interested parties;		
<ul><li>2) the extent to which quality objectives have been met;</li></ul>		
<ul> <li>a) process performance and conformity of products and services;</li> </ul>		
<ul> <li>a) process performance and connormity of products and services,</li> <li>a) nonconformities and corrective actions;</li> </ul>		
5) monitoring and measurement results;		
6) audit results;		
7) the performance of external providers;		
d) the adequacy of resources;		
e) the effectiveness of actions taken to address risks and opportunities (see <u>6.1</u> ); f) opportunities for improvement.		
9.3.3 Management review outputs The outputs of the management review shall include decisions and actions related to:		
<ul> <li>a) opportunities for improvement;</li> </ul>		
<li>b) any need for changes to the quality management system;</li>		
c) resource needs.		
The organization shall retain documented information as evidence of the results of management reviews.		
10 Improvement		
10.1 General		
The organization shall determine and select opportunities for improvement and implement any necessary actions to		
meet customer requirements and enhance customer satisfaction.		
These shall include:		
a) improving products and services to meet requirements as well as to address future needs and expectations;		
b) correcting, preventing or reducing undesired effects;		
	-	•



c) improving the performance and effectiveness of the quality management system.	
NOTE Examples of improvement can include correction, corrective action, continual improvement,	
breakthrough change, innovation and re-organization.	
10.2 Nonconformity and corrective action	
<b>10.2.1</b> When a nonconformity occurs, including any arising from complaints, the organization shall:	
a) react to the nonconformity and, as applicable:	
1) take action to control and correct it;	
2) deal with the consequences;	
b) evaluate the need for action to eliminate the cause(s) of the nonconformity, in order that it does not recur or occur	
elsewhere, by:	
<ol> <li>reviewing and analysing the nonconformity;</li> </ol>	
2) determining the causes of the nonconformity;	
<ol> <li>determining if similar nonconformities exist, or could potentially occur;</li> </ol>	
c) implement any action needed;	
d) review the effectiveness of any corrective action taken;	
e) update risks and opportunities determined during planning, if necessary; f) make changes to the quality	
management system, if necessary.	
Corrective actions shall be appropriate to the effects of the nonconformities encountered.	
<b>10.2.2</b> The organization shall retain documented information as evidence of:	
a) the nature of the nonconformities and any subsequent actions taken;	
10.3 Continual improvement	
The organization shall continually improve the suitability, adequacy and effectiveness of the quality management system.	
The organization shall consider the results of analysis and evaluation, and the outputs from management review, to	
determine if there are needs or opportunities that shall be addressed as part of continual improvement.	
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