

9100 AEROSPACE INDUSTRY SPECIFIC COURSE CHECKLIST

(Formerly the in-depth/competency course.)

	LEARNING OBJECTIVES	OBJECTIVE STATED/COVERED IN COURSE PLAN		
Α	Explain the quality perspective of the aerospace industry.	☐ YES	□ NO	
В	Describe in overview CAA roles, responsibilities, and regulations, including how to obtain additional information.	☐ YES	□ NO	
С	Describe the principles of military aerospace requirements and regulations, including how to obtain additional information.	☐ YES	□ NO	
D	Describe in overview government space organization roles, responsibilities, and regulations, including how to obtain additional information.	☐ YES	□ NO	
Е	Describe in overview airworthiness and aviation safety requirements, including how to obtain additional information.	☐ YES	□ NO	
F	Describe the design, development, verification, and validation processes specific to the aerospace industry.	☐ YES	□ NO	
G	Describe control of production and service provision.	☐ YES	□ NO	
Н	Explain the process of first article inspection (see 9102).	☐ YES	□ NO	
ı	Describe aerospace material traceability requirements.	☐ YES	□ NO	
J	Describe aerospace material/parts status accountability systems.	☐ YES	□ NO	
K	Identify aerospace subcontractor approval and control requirements.	☐ YES	□ NO	
L	Explain variation management of key characteristics (see 9103).	☐ YES	□ NO	
М	Describe flow down of quality requirements.	☐ YES	□ NO	
N	Explain FOD prevention program requirements.	☐ YES	□ NO	
0	Describe use of customer supplied products.	☐ YES	□ NO	
Р	Explain calibration controls and positive recall system.	☐ YES	□ NO	
Q	Describe acceptance authority media (i.e., use of stamps, stamp control, electronic sign off).	☐ YES	□ NO	
R	Identify special processes.	☐ YES	□ NO	
S	Identify nonconforming material, system requirements, and operation.	☐ YES	□ NO	
Т	Explain sampling inspection requirements and limitations.	☐ YES	□ NO	
U	Identify configuration management/requirements control.	☐ YES	□ NO	
V	Describe tool control.	☐ YES	□ NO	
W	Describe the product qualification process.	☐ YES	□ NO	



Χ	Describe/explain source approval process.	☐ YES	□ NO				
	aviation authorities and regulations covered in the course shall be applicable to the region in the course is being presented.	☐ YES	□ NO				
	30% (8 hours) of the total course time shall be used for active participation in workshop, case studies, role-playing, and/or actual auditing of an organization as part of the structured class activities.						
	ESTIMATED TIME: ACTUAL TIME:						
	The total course time devoted to direct instruction and to assigned team and individual activities shall be at least 26 hours , plus an additional hour for the written examination.						
	ESTIMATED COURSE TIME: ESTIMATED EXAM TIME: _						
EVALU	ATOR NOTES:						



NO.	TRAINING ITEM / LEARNING OBJECTIVE	TRAINING SUBJECTS	MET	ESTIMATED HOURS	ACTUAL HOURS
1	Aerospace industry quality perspective.	History of the standard, reasons for introduction, relation with airworthiness requirements.		1	
2	CAA requirements.	Difference between customer quality requirements and airworthiness requirements. Major content of EASA Part 21 and part 145, relation to FAR requirements, Design Organization Approval (DOA), Production Organization Approval (POA), and Manufacturing Organization Approval (MOA) approvals; type certification, supplier control principles, certifying staff, accountable manager, and use of EASA Form 1.		2.5	
3	Principles of military aerospace requirements and regulations.	Use of Allied Quality Assurance Publication (AQAP), status, military as customers, relation to contractual requirements.		1	
4	Principles of space requirements and regulations.	Government space organization roles, responsibilities, and standards.		1	
5	First Article Inspection (FAI) (see 102).	Purpose of FAI, structure and use of 9102, and supporting forms.		2	
6	Airworthiness and safety requirements.	See CAA requirements, difference and relation between safety and quality; responsibility of supplier and customer; and position of OEM and subcontractor.		Included in No. 2	
7	Design, development, verification, and validation.	The management model for design, design and change control, difference between verification and validation; type certification and approved design data.		2	
8	Aerospace material traceability requirements.	Upwards and downwards traceability, lot control, serialized items, marking, records for traceability, and traceability of raw material.		1	
9	Aerospace material accountability system.	Accounting for all parts during manufacturing (e.g., split batches).		0.5	



10	Aerospace subcontractor approval and control requirements.	Responsibility for quality, supplier auditing, Approved Supplier List, function of incoming and source inspection.	2	
11	Variation management of key characteristics (see 9103).	Principles of statistical control, attributive and contributive measuring, Cp and Cpk, 6 Sigma, method(s) and approach to reduce variation, and criteria to establish key characteristics.	2	
12	Flow down of quality requirements.	Sub-tier control, role of the contract, and purchasing data.	Included in No. 10	
13	Foreign Object Damage/Debris (FOD) prevention program requirements.	Definition of FOD, FOD prevention, FOD inspections/detections, and FOD audits.	1	
14	Use of customer supplied products.	Responsibility for quality of customer supplied product and use of customer approved sources.	0.5	
15	Calibration controls and positive recall system for monitoring and measuring devices.	Description of 'positive' recall, administration, use of stickers, release after calibration, sub- contracting of calibration, and actions in case of 'out of calibration'.	1	
16	Acceptance authority media.	Use of stamps, stamp control, marking, and electronic sign off.	1	
17	Nonconforming material system requirements and operation.	Responsibilities, identification, quarantine rules, Material Review Board, dispositions, release of nonconforming products, and records.	2	
18	Sampling inspection requirements and limitations.	Use of sampling techniques and sampling standards.	1	
19	Tool control.	Control of tool including traceability, 'positive' recall, administration, use of stickers, and release after calibration.	0.5	
20	Special processes.	Definition of special processes, examples of special processes, special process qualification, control, and special process audits.	2	



21	Configuration management/requirements control.	Definition/description of configuration management, configuration management policy, Configuration Items (CI), configuration control, configuration auditing, relation with traceability requirements, and configuration changes.	2	
			26	