

Applicant Name:

European Aviation Safety Agency

Form

Tel No:

Part-21 SUBPART G PRODUCTION ORGANISATION EXPOSITION COMPLIANCE CHECKLIST

Contact Person:			Approval Ref:	
POE Title:			POE Ref:	
Date of Review:			Reviewed by:	
			,	
This is the list of all the chapter the chapters but it is mandatory Some chapters can be added acc The titles can be changed if app	to cover all the cording to Organi	ones applicable.	-	(POE). It is not mandatory to follow the sequence of
	Reference to EASA Part 21 subpart G	Ref to POE paragraph	Comment for applicant	Comment for Competent Authority
General information that should be in the first page				
Part 21 subpart G Production Organisation Exposition				
Name and address of the Organisation complying with official name (EF50 and business registration)				
Approval reference of the POA				
Reference of the Exposition with issue number				
Approval date				
General information for each page]			
Name of the organisation				
POE identification				
Amendment/revision number of the POE				
Page number				
General chapters				
Table of content				

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History of revision	Including status of the revision. Please ensure that the changes are somehow highlighted and that they are easy to identify.	
List of effectives pages		
Distribution list		
Terms and abbreviation	This can be removed from general chapters if any abbreviations is defined every time it is used in the document	
Introduction / Description of the Organisation	This is to present the organisation	

Management Procedures			
Signed corporate commitment by the Accountable Manager	21.A.143 (a) 1.		
Nomination of Accountable Manager with reference to delegation letter when the AM is nominated by top management	21.A.143 (a) 2. 21.A.145 (c) .1	Shall confirm that the production organisation exposition and any associated manuals which define the approved organisation's compliance with this subpart will be complied with at all times.	
Management personnel	21.A.145 (c) 2. 21.A.143 (a)	Shall list the title and names of all the nominated persons in front of the POA with identification of EASA Form 4 holders	
Duties and responsibilities of :	21.A.143 (a) 3. 21.A.145 (c) 2.	Shall also include matters on which they may deal directly with the competent authority on behalf of the Organisation.	
- Accountable manager			
- Quality manager			
- Production manager			
 Any other manager related to POA 			
Organisation chart	21.A.143 (a) 4. 21.A.145	The org chart shall identify the reporting lines and nominated managers	
List of Part 21 certifying staff	21.A.143 (a) 5. 21.A.145 (d)	This can also be an appendix	
General description of the man-power resources	21.A.143 (a) 6.		
General description of the facilities	21.A.143 (a) 7.	Containing the address and details of each facility included in the scope of the POA (in the production organisation's certificate of approval). A readable facility layout plan shall be included	
Scope of work	21.A.143 (a) 8. 21.A.151	The general scope of work relevant to the terms of approval shall be described here.	

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		Additionally it should refer to the full list of P/N (part number) produced under the production approval, the capability list or to the database that gives the list. For the products, it should refer to the type certificate number. In case of various DO/PO arrangements, a list of all DO/PO arrangements shall be included.	
Notification procedure of organisational changes to Competent Authority.	21.A.143 (a) 9. 21.A.147 (a) 21.A.148 21.A.149 21.A.153	Shall list all the changes identified as significant changes. Shall describe how each type (significant or not) of changes are managed. It includes change of accountable manager, change of other nominated managers, change of location of facility or change of activity (scope) etc	
Amendment procedure of the exposition	21.A.143 (a) 10. 21.A.143 (b) 21.A.165 (a)	It shall describe how and by whom are the Exposition and the associated documents updated.	
Description of the quality system	21.A.143 (a) 11.	This is optional as it is covered by the next chapter but it can be useful to describe the structure of the documentation (pyramid)	
Supplier/subcontractor list	21.A.143 (a) 12	It shall include the main suppliers list plus the reference to the full suppliers list if the list is too big. A change of such a main subcontractor may be treated as a significant change (21.A.147 (a)). Can also be put as an appendix.	

Quality System]		
Distribution of the documents	21.A.139 (a) 21.A.165 (a)		
Document issue, approval or change	21.A.139 (b) 1. (i)	The creation of document (by whom, to whom, numbering, document structure) shall also be covered in this paragraph. How the changes are followed and highlighted shall also be covered.	
Vendor and subcontractor assessment audit and control	21.A.139 (b) 1. (ii) 21.A.157	Shall also include the evaluation and the acceptance criteria.	
Verification that incoming products, parts, materials, and equipment, including items supplied new or used by buyers of products, are as specified in the applicable design data	21.A.139 (b) 1. (iii)	It is the description of the incoming material inspection	
Identification and traceability	21.A.139 (b) 1. (iv)		



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Manufacturing processes	21.A.139 (b) 1. (v) 21.A.145 (a) 21.A.163 (a) 21.A.165 (b)	Shall also include the management of the production documentation.	
Special processes	21.A.145 (d)	The special processes shall be mentioned and described if any.	
Inspection and testing, including production flight tests	21.A.139 (b) 1. (vi)		
Calibration of tools, jigs and test equipment	21.A.139 (b) 1. (vii)	Shall include the acceptance, the use, the control and the calibration of the tools and equipment	
Non-conforming items control	21.A.139 (b) 1. (viii)	Including concessions	
Airworthiness co-ordination with applicant for, or holder of, the design approval	21.A.139 (b) 1. (ix) 21.A.133 (b) (c) 21.A.165 (g)	This paragraph shall also refer to the DO/PO arrangement if any (unless this is included in the "scope of work" chapter).	
Records completion and retention	21.A.139 (b) 1. (x) 21.A.165 (d) 21.A.165 (h)	It is dealing with technical records and it shall include the management of electronic records if any.	
Personnel competence and qualification	21.A.139 (b) 1. (xi) 21.A.145 (d)	This should describe the general requirement for accepting anybody working in POA holder organisation. The training process of these persons shall be described (minimum training and also regular training). If there are special process or NDT in the scope, the specific requirements for training and qualification should also be described.	
Certifying staff qualification and training	21.A.145 (d)	This paragraph is specifically reserved for certifying staff, with qualification requirements, training needs, nomination, records and authorization.	
Issue of airworthiness release documents	21.A.139 (b) 1. (xii) 21.A.163 21.A.165 (c) 21.A.165 (i)		
Handling, storage and packing	21.A.139 (b) 1. (xiii)		
Internal quality audits and resulting corrective actions	21.A.139 (b) 1. (xiv) 21.A.139 (b) 2. 21.A.158		
 Quality audit of processes Quality audit of product Quality audit remedial action procedure Quality audit personnel Planning for POA compliance audits 		The quality audit of processes shall cover also the audit of special processes if any. These are the audits procedures to cover the scope of Part 21 subpart G in order to prove the compliance with the regulation	

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Work within the terms of approval performed at any location other than the approved facilities		21.A.139 (b) 1. (xv)	Also called outlocated work.		
Work carried out after completion of production but prior to delivery, to maintain the aircraft in a condition for safe operation		21.A.139 (b) 1. (xvi)	This is applicable only for complete aircraft.		
Issue of permit to fly and approval of associated flight conditions		21.A.139 (b) 1. (xvii) 21.A.165 (j) (k)	This is applicable only for complete aircraft.		
Occurrence reporting		21.A.139 (f) 21.A.165 (e) (f)			
Control of critical parts		21.A.139 (b) 1.			
Appendi	ixes	7			
Capability List			If applicable		
Cross reference table between Part 21 subpart G requirements and internal			This is not applicable in case there are no other internal POA documents than POE.		

Date:

Conclusion/Notes:

documents.

Reviewed by:

Signed:



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