IATF - International Automotive Task Force Rules for achieving and maintaining IATF Recognition, 6th Edition --- Sanctioned Interpretations (SIs)



IATF - International Automotive Task Force **Rules for achieving and maintaining IATF Recognition IATF Rules6th Edition – Sanctioned Interpretations**

The *Rules for achieving and maintaining IATF Recognition 6th Edition for IATF 16949* ("**Rules 6th Edition**") was published in April 2024 and is effective 1 January 2025. The following Sanctioned Interpretations were determined and approved by the IATF. Unless otherwise indicated, Sanctioned Interpretations are applicable upon publication.

Revised text is shown in **blue**.

A Sanctioned Interpretation changes the interpretation of a rule or a requirement which itself then becomes the basis for a nonconformity.

- SI 1 SI 17 issued November 2024, effective 1 January 2025
- SI 7 revised and reissued April 2025, effective April 2025
- SI 18 issued April 2025, effective April 2025

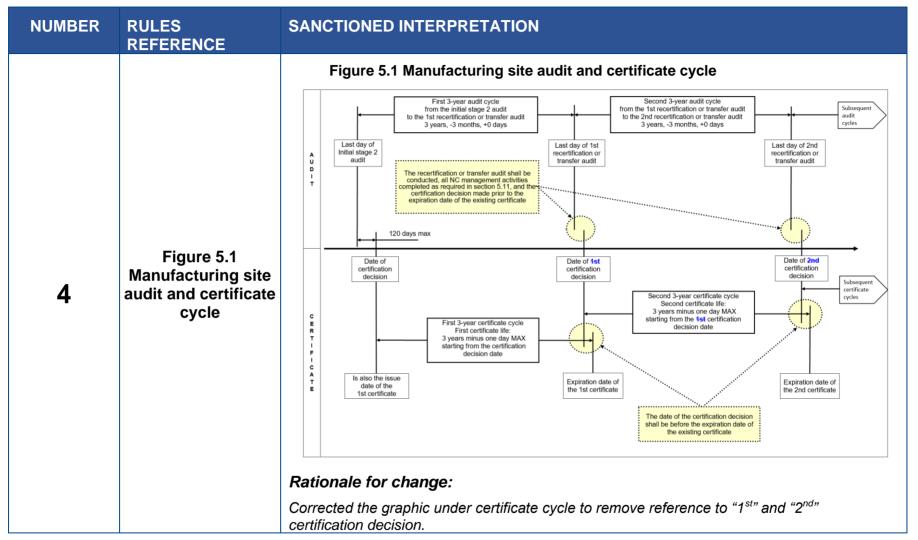


NUMBER	RULES REFERENCE	SANCTIONED INTERPRETATION
1	Foreword	Requests for permission to reproduce and/or translate this document or any extract from it must be addressed to one of the organizations below: Automotive Industry Action Group (AIAG/USA) Associazione Nazionale Filiera Industria Automobilistica (ANFIA/Italy) International Automotive Task Force France (IATF France/France) IATF France (France) The Society of Motor Manufacturers and Traders Limited (SMMT/UK) Verband der Automobilindustrie e.V. – Qualitäts Management Center (VDA QMC/Germany) Rationale for change: Clarified the names of the copyright owners in France, UK, and Germany.
2	Global overview of the first 3-year IATF 16949 certification cycle	Global overview of the first 3-year IATF 16949 certification cycle Plan for stage 2 Conduct (5.7.1 & 5.7.2) Plan for stage 2 0 days and no more than 90 days after Plan for surveillance audit (5.7.1 & 5.7.2) Plan for recertification audit of remote Somer than support locations Plan for surveillance where applicable Conduct stage 2 (5.5.1 - 5.5.2) Conduct surveillance Conduct stage 2 Conduct surveillance audit (5.8 & 6.2.9) Conduct surveillance Issue draft audit report at closing meeting (5.10) Issue draft audit report at closing meeting (5.10) Rationale for change: Corrected reference to 6.3 (was 6.7) in the "Conduct surveillance audit" step.



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3		SANCHONED INTERFECTIONTable 2.10 Minimum number of witness audits to perform during the interf varia during the interf varia during the 						
		Removed the word "Note" as the increase of one (1) witness audit per every 1,500 audit days after 31,000 audit days is a requirement not guidance.						







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5	5.2 Determining the audit duration for initial certification, surveillance, recertification, and transfer audits	 When determining the audit duration and developing an audit plan (see section 5.7.2), the certification body shall consider, at minimum, the following: a) - p) all other requirements of section 5.2 remain unchanged q) When the audited client location does not meet the IATF OEM quality and/or delivery targets specified in the IATF OEM scorecard(s) submitted with audit planning information and/or the current scorecards reviewed in the opening meeting, the certification body shall add "additional audit time" to the current audit plan as defined in Table 5.2 q. The "additional audit time" shall be used to verify the effective implementation of the systemic corrective actions associated with the IATF OEM quality and/or delivery targets that are not being met and the associated risk to similar processes and/or products. The only exception is if the client can provide evidence of verification of the effectiveness of the implemented systemic corrective actions for the quality and/or delivery performance issues, in which case no increase is applied. Where it is not possible to add the additional time to the current audit plan, the certification body shall conduct a special audit within sixty (60) calendar days of the closing meeting date of the audit (see section 7.2 c) to 7.2 b). 			



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6	Minimum audit days Table 5.2	

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		The certification body shall have a process for audit planning in accordance with the relevant requirements provided in ISO/IEC 17021-1 and the basic requirements stated below.
		Audit planning shall be completed before the start of every audit.
		Note: Stage 1 readiness assessment planning requirements can be found in section 6.2.1 of these Rules.
		Each manufacturing site, including those in a corporate scheme, and any standalone remote support location shall have dedicated audit planning for each audit.
		Extended manufacturing sites shall be included in the audit planning for the main manufacturing site.
7	5.7	The certification body shall provide for a minimum of one-half (0.5) day for audit preparation and planning for each manufacturing site. The certification body shall determine the minimum audit preparation and planning time for each stand-alone remote support location audit and special audit based on risk and audit scope. The audit preparation and planning time shall be entered into the IATF Database. ² This time shall not be included in the audit duration (see section 5.2).
(Revised)	Audit planning	A member of the audit team shall undertake an analysis of the audit planning information provided by the client and the most recent audit records from relevant remote support locations (see section 5.5.3) to determine critical areas to be prioritized for the audit based on risk to the customer, performance trends, and criticality of the client's processes. The analysis may result in an adjustment to the audit duration. Justification for not adding "additional audit time" to investigate IATF OEM customer quality and delivery performance issues (see section 5.2 q) shall be recorded. Analysis results shall be recorded to include the minimum content in Annex 3 – Table for Documenting the Output of the Planning Process and retained as part of the audit records.
		For stage 2 certification audits, audit planning shall consider facts established during the stage 1 readiness assessment.
		For recertification audits, audit planning shall include a review of internal and external performance data and nonconformities reported in surveillance audit reports from the current audit cycle, as needed, to evaluate the performance of the quality management system during the period of the current certification and to identify any areas that need to be prioritized during the audit.
		A member of the audit team shall review the client's website to validate the audit planning information provided by the client (e.g., organization structure, customers, support activities, the

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		certification scope, use of the IATF logo, etc.).			
		Audit start and end ¹ dates for surveillance, recertification, and transfer audits shall be confirmed with the client no less than ninety (90) calendar days before the audit due date (see section 10.0). If an exceptional situation requires a change to a confirmed audit date, the certification body shall retain justification as a part of the audit record.			
		The certification body and the client shall plan the audit to ensure that automotive manufacturing processes will be running as required during the planned audit duration (see section 5.8.5). If this requirement cannot be met, the certification body shall delay the audit until the requirement can be met.			
		Note: Delaying an audit may result in loss of certification.			
		Rationale for change:			
		¹ To clarify what dates must be confirmed ninety calendar days before the audit due date.			
		² To allow the CB to determine the appropriate amount of time for audit preparation and planning time for SA-RSLs.			
		The audit team shall analyze all information and objective evidence gathered during the audit and agree on the audit conclusion.			
	5.10 Audit reporting	The audit team shall use the IATF Common Audit Report Application (IATF CARA) when creating the draft and final audit reports and for the issuance of nonconformities to the client. All mandatory fields within the IATF CARA shall be completed with sufficient detail and objective evidence to allow the reader to comprehend the audit coverage, findings, results, and conclusions.			
8		The certification body shall issue the draft audit report and the nonconformity management record(s), where applicable, to the client at the audit closing meeting along with the link to the IATF Common Audit Report Application for Nonconformity Management (IATF NC CARA).			
		The remainder of section 5.10 remains unchanged			
		Rationale for change:			
		To clarify the term "nonconformity management record" includes the client's response and the term "nonconformity record" is just the CB auditor's finding and to clarify the IATF NC CARA link is to be provided at the closing meeting.			



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9	5.11.4 Verification of major nonconformity	In cases of a major nonconformity, the certification body shall conduct a one-time, onsite special audit (see section 7.2 d c]) for the verification of the effective implementation of systemic corrective actions within a maximum of ninety (90) calendar days from the closing meeting date of the audit. The remainder of section 5.11.4 remains unchanged. <i>Rationale for change: Corrected the cross reference from section 7.2 d) to 7.2 c)</i>			
10	5.12 Technical review and certification decision	The certification decision shall be made within a maximum of one-hundred-twenty (120) calendar days from the last day of a special audit (see section 7.2 e), f), g) and h)) unless the special audit is being conducted within the decertification process timing. The remainder of section 5.12 remains unchanged. Rationale for change: Added clarification to which type of special audits requires a certification decision within one- hundred-twenty (120) calendar days.			
11	6.1.1 Application for certification	 The certification body shall require a representative of the applicant organization to provide the necessary information to enable the certification body to decide on certification feasibility and to establish a complete quotation. The information required shall include, at minimum: a) - n)these requirements remain unchanged o) All IATF 16949 certifications within the last three (3) years, the status of the certificate(s), and all final audit reports and all nonconformity management records from the last three (3) years of certification, including evidence that all nonconformity responses have been accepted and all verification activities that should have been completed by the previous certification body were conducted as required (see section 5.11). If the certification body is unable to determine the reason for the applicant organization's certification withdrawal from the information on the reason for the withdrawal. p) - r) all other requirements remain unchanged Rationale for change: To clarify the term "nonconformity management record" includes the client's response and the			

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		term "nonconformity record" is just the CB auditor's finding.			
		If a client chooses to change to another IATF-recognized certification body, the location(s) for which the transfer is requested shall undergo a transfer audit with the new certification body.			
		Clients may apply for transfer at any point in the audit cycle.			
		The transfer audit shall occur within the allowable timing for the next regular audit with the previous certification body (see sections 5.1.1 and 5.5.1).			
	7.1 Transfer audit	Within seven (7) calendar days of a legal contract being signed with the client, the new certification body shall indicate the transfer in the IATF Database by creating the client record and completing the information required in the transfer confirmation feature.			
		The new certification body shall advise the client to notify the previous certification body about its intent to transfer (see section $3.2 \vdash j$).			
12		The previous certification body shall not use the transfer intent notification as justification for changing the client's certificate status before the transfer process is complete, provided that no other valid justification exists, and the client and the previous certification body still have a valid contract in place.			
		Unless otherwise explicitly stated in these Rules, the certification activities for a transfer audit shall be the equivalent of a recertification audit in terms of audit scope, duration, planning, and conduct, and the relevant requirements for recertification audits apply (see section 6.4).			
		The new certification body shall only conduct an initial certification audit (i.e., stage 1 readiness assessment and stage 2 certification audit) instead of a transfer audit if any of the following conditions exist:			
		The transferring manufacturing site does not have a valid certificate or only has a letter of conformance.			
		The transferring standalone remote support location is not referenced on a valid manufacturing site certificate or only has a letter of conformance.			
		The allowable audit timing for the transfer audit has been exceeded.			
		Note: These Rules may require the new certification body to conduct special audits before the start of the initial certification audit (see sections 6.1.1, 7.2, and 8.7).			



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		The certification body shall enter all the required audit details in the IATF Database within twenty (20) calendar days from the closing meeting of the transfer audit. This information shall be in the specified format and in English.			
		Rationale for change:			
		Corrected the cross reference from 3.2 i) to 3.2 j).			
		7.1.2 7.1.1 Transfer audit pre-conditions			
		At the start of the transfer audit, the following conditions shall be met for the client location under transfer:			
	7.1.2 Transfer pre- conditions	a) – h) …these requirements remain unchanged.			
13		 i) The client provided the new certification body with the final audit reports and nonconformity management records from the previous three (3) years. The information provided shall include evidence that client responses to all previously issued nonconformities have been accepted and all verification activities that should have been completed by the previous certification body were conducted as required (see section 5.11). 			
		Wherever possible, open issues from the points above shall be resolved by the previous certification body and the client.			
		Rationale for change:			
		To correct the section header number from 7.1.2 to 7.1.1. To clarify the term "nonconformity management record" includes the client's response and the term "nonconformity record" is just the CB auditor's finding.			



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14	10.0 Definitions	 Certification body (IATF-recognized) The Named Certification Body, together with all Affiliates that satisfy the conditions required to conduct IATF Certification Activities pursuant to sections 1.6.1 and 1.6.2 in the "Agreement". Certificate scope statement The statement displayed on the IATF 16949 certificate or the letter of conformance that indicates reflects the manufacturing processes and, where applicable, product design responsibility, covered the products being designed and/or manufactured under the quality management system. The certificate scope statement only reflects may also include the categories of automotive products the client manufactures. is manufacturing at the site that is being certified, and, where applicable, indicates the client's responsibility for their design. The certificate scope statement may also include the client's manufacturing processes and/or technologies used to further convey the character of the automotive products. Consecutive working days Calendar days Regular working days of the client over which an audit is conducted in succession and without interruption. Rationale for change: Align the words in section 10.0 with terms used in section 5.7 and 5.7.2 related to client working days, section 5.13 for certificate scope statement and to remove a reference in the sections in the CB Agreement that changed.



	RULES REFERENCE	SANCTIO	SANCTIONED INTERPRETATION						
		Example 2 1 single manufac 1 permitted redu No standalone re	Annex 1 – Example 2 & 4 Example 2 1 single manufacturing site with 1 extended manufacturing site (main site: 550 employees, EMS: 40 employees) 1 permitted reduction (no design responsibility) No standalone remote support locations Total number of employees = 590 (550 + 40)						
					Current number of				Minimum audit day requirement (rounded
		Site	Year	Type of audit	employees		reductio		up to nearest 1/2 day)
		1 (main + EMS)	0	Initial Stage 2	590	10.5	15%	(10.5 days * 0.85) = 8.925	9.0
		1 (main + EMS)	1	Surveillance	590	5.25	15%	(5.25 days * 0.85) = 4.4625	4.5
		1 (main + EMS) 1 (main + EMS)	2	Surveillance Recertification	590 590	5.25	15% 15%	(5.25 days * 0.85) = 4.4625 (7.0 days * 0.85) = 5.95	4.5
15	Example 2 Example 4	Total number of employees = 590 (550 + 40) Using the apportionment method							
		Site	Year	Type of audit			Non-o grade respo n LOC redu	nsible	Minimum audit day requirement (rounded up to nearest 1/2 day)
			Year 0	Type of audit	number of	day requirement, Up Rules Table 5.2 fro	grade respo	ction Calculated minimum audit d	requirement (rounded up to lays nearest 1/2 day)
		Site 1 (Main + EMS) 1 (Main + EMS)	0		number of employees 590 590	day requirement, Up Rules Table 5.2 from 10.5 5.25	respondent of the second secon	calculated minimum audit d % (10.5 days * 0.70) = 7.35 % (5.25 days * 0.85) = 4.462	requirement (rounded up to nearest 1/2 day) 5 7.5 25 4.5
		Site 1 (Main + EMS)	0	Initial Stage 2	number of employees 590	day requirement, Up Rules Table 5.2 from 10.5	n LOC response response redu	calculated minimum audit d % (10.5 days * 0.70) = 7.35 % (5.25 days * 0.85) = 4.462 % (5.25 days * 0.85) = 4.462	requirement (rounded up to nearest 1/2 day) 5 7.5 25 4.5 25 4.5



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16	Annex 1 Example 3	Annex 1 - Example 3 Simulation of the support line of the stretched many functions (main sket. 550 employees, EMS: 40 e								



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BER RULES SANCTIONED INTERPRETATION REFERENCE	
Annex 1 – Example 7 Corporate certification structure 3 single manufacturing sites (site 1: 590 employees, site 2: 800 employees, site 3: 850 employees) 2 semilted reductions, including a central location (Total of 125 employees) 2 permitted reductions (non-design responsibility and corporate certification structure) Using the apportionment method Number of employees at site / total Number of employees at site / total Site Total number of employees at site / total Site Total number of employees at site / total 1 Site Site Total number of employees at site / total 1 Site Total number of employees at site / total 1 Site Total number of employees at site / total 1 590 26% 32 622 2 800 36% 48 898 Total = 125 Total = 2,365	
Site Number of employees at site Minimum audit day requirement, Rules Table 5.2 Non-design responsible reduction Corporate catification structure reduction Calculated mininum audit days day (re mea 1 (Main & EMS) 622 11.0 15% 15% (11.0 days * 0.70) = 7.7	Ainimum audit ay requirement rounded up to arest 1/2 day) 8.0
Annex 1 Example 7	8.5 8.5
Site Number of employees at requirement, Rules reduction days (if days in the formation of the section days in the section days (if	Ainimum audit ay requirement rounded up to earest 1/2 day)
1 (Main & EMS) 622 5.5 15% 15% (5.5 days * 0.70) = 3.85	4.0
2 845 6.0 15% 15% (6.0 days * 0.70) = 4.2 3 898 6.0 15% 15% (6.0 days * 0.70) = 4.2	4.5 4.5
Total = 2,365 Calculation for Recertification Audit: Minimum audit day requirement, Rules Non design responsible reduction Corporate certification days Minimum audit day requirement, Rules	finimum audit ay requirement rounded up to arest 1/2 day)
1 (Main & EMS) 622 7.0 15% (7.0 days * 0.70) = 4.9	5.0
2 845 7.5 15% 15% (7.5 days * 0.70) = 5.25	5.5
3 998 8.0 15% (8.0 days * 0.70) = 5.6	6.0
Total = 2,365 Note 1: Assumes no changes over the 3-year audit cycle to number of sites or SA-RSLs, number of employees, certification scope, etc. It is the responsibility of the certification body to determine the distribution of the minimum audit days each year from the manufacturing sites to the remote support locations to ensure the effectiveness of the audits.	he standalone
Rationale for change:	
Corrected to remove the main site + EMS from Site 1 to align with	

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IO red		When creating an audit program for manufacturing sites with support functions performed onsite, the certification body shall meet the following requirements:
		a) Product design functions shall be included in the initial stage 2 audit, in each surveillance audit, and in the recertification audit. All other support functions shall be included in the initial stage 2 audit, at least once more during the surveillance audit cycle, and in the recertification audit.
		b) Stage 2 certification audit planning for a manufacturing site shall ensure that all its support functions performed at standalone remote support locations are audited and relevant interfaces validated at the standalone remote support locations before the audit at the manufacturing site.
		When creating an audit program for standalone remote support locations, the certification body shall meet the following requirements:
	5.5.1 Audit program	c) The audit program shall start with an initial certification audit—or a transfer audit, where applicable—followed by an ongoing sequence of surveillance audits.
	requirements for support functions	 Audits shall be programmed and conducted based on audit due dates established from the initial certification audit or transfer audit date.
		 Product design functions shall be audited annually (i.e., every twelve [12] months [-3/+3 months]).
		 f) Central locations within a corporate scheme shall be audited annually (i.e., every twelve [12] months [-3/+3 months]).
		g) Non-product design functions shall be audited at least every two (2) years (i.e., every twenty- four [24] months [-3/+3 months]).
		 Failure to conduct audits of the standalone remote support location within the required timing shall result in an initial certification audit.
		Rationale for change:
		Clarified the central location within a corporate scheme is required to have an annual audit.

