



International Management Accreditation Board

Address : 51, Goldhill Plaza, #07-10/11, Singapore 308900
 E-mail : info@imacb.com, customersupport@imacb.com
 Website : www.imacb.com

Necessary Document Checklist
 for Accreditation of Inspection Bodies in accordance with DIN EN ISO/IEC 17020:2012

Name of Inspection Body	
Address of Inspection Body	
Type of Inspection Body	
Case number	
Date of document submission	

The required documents shall preferably be submitted electronically, in a way that the numbering can directly be assigned to the relevant documents. Please send fully updated documents to customer manager. In individual cases, documents may be submitted in hard copy, the customer manager will inform you if necessary.

All documents/records shall be submitted in a timely manner for each assessment. If necessary, further documents may be requested by the customer manager or by assessors.

No.	Document	If applicable cross reference to QM	Ok
1.	Quality manual (QM)		<input type="checkbox"/>
2.	Master list(s) of all QM documents		<input type="checkbox"/>
3.	Services provided by the inspection body		<input type="checkbox"/>
4.	Proof of organisation, ownership and legal entity (e. g. extract from a registered statement)		<input type="checkbox"/>
5.	Proof of third party liability insurance		<input type="checkbox"/>
6.	Independence declaration of the top management according to type A, B, C		<input type="checkbox"/>
7.	Evaluation of the risk of impartiality (Analysis of related bodies)		<input type="checkbox"/>
8.	Staff declaration of confidentiality or evidence of such		<input type="checkbox"/>
9.	Organisational chart		<input type="checkbox"/>
10.	List of employees stating their qualification/professional training		<input type="checkbox"/>



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11.	Evidence of qualification of the head of the inspection body and his/her deputy		<input type="checkbox"/>
12.	List of staff members who are approved signatories of inspection reports stating their evaluation competence and specimen signatures		<input type="checkbox"/>
13.	Evidence of qualification of the staff members who are approved signatories		<input type="checkbox"/>
14.	Copy of at least one original inspection report for each inspection area scheduled for accreditation		<input type="checkbox"/>
15.	Documentation of inspection cases for each scheduled inspection program / each scheduled area of inspection		<input type="checkbox"/>
16.	If applicable, documentation of in-house methods of tests and their validation		<input type="checkbox"/>
17.	If applicable, list of reference materials in use		<input type="checkbox"/>
18.	Information on the participation in proficiency tests such as interlaboratory comparisons stating the date, organizer, parameters, matrix, results and conclusions in tabular form		<input type="checkbox"/>
19.	List of equipment items with in-house registry (including loaned equipment, if applicable) Required information: Inventory number, location, indication of the equipment/ type of equipment/item, manufacturer, calibration interval, date of the next calibration, indication of the proof of measurement traceability, kind of calibration certificate. Optional information : Testing standard, serial number, responsible person for the equipment, etc.		<input type="checkbox"/>
20.	Site plan showing the work areas		<input type="checkbox"/>
21.	If necessary and desired: Approval/withdrawal of approval for forwarding documents and data to third parties (please specify)		<input type="checkbox"/>
22.	Requirements for handling the above mentioned documents/records, when outdated or replaced by others		<input type="checkbox"/>