**Recognition Requirements for Conformity Assessment Bodies**

This document outlines the IMDRF GRRP recognition requirements for Conformity Assessment Bodies conducting medical device regulatory reviews and is intended to aid in the review of the draft IMDRF document entitled *Regulatory Authority Assessment Method for Recognition and Surveillance of Medical Device Conformity Assessment Bodies*. ISO/IEC 17065:2012 is being utilized as a normative reference for the requirements outlined below and the clause numbers below are aligned with the relevant subsection in ISO/IEC 17065:2012. Specifically, Sections 4-8 align with the requirements of the standard and additional requirements are included in Sections 9-10. “Exceptions to ISO/IEC 17065:2012” listed below mean that certain requirements in ISO/IEC 17065:2012 do not apply to marketing review conformity assessment. “Specific requirements for medical device CABs” are requirements for marketing review in addition to those outlined in the corresponding section of ISO/IEC 17065:2012.

# General Requirements

## Legal and Contractual Matters

Country-specific laws and regulations, outside the medical device Regulatory Authority’s purview, may be applicable to the manufacturer for certain legal and financial responsibilities.

A legal entity is ineligible to be a medical device CAB if the entity has been found guilty of an offense against national laws or regulations related to medical devices, or relating to any fraudulent or dishonest practices.

**Exceptions to ISO/IEC 17065:2012**

In cases where the Regulatory Authority issues the final marketing decision, the agreement between the CAB and the manufacturer shall require the manufacturer to notify the Regulatory Authority of changes affecting conformity with the certification decision discussed in 4.1.2.2(k).

**Specific Requirements for Medical Device CABs**

### A CAB shall make available to the recognizing Regulatory Authority(s) information about organizational structure, ownership and the legal or natural persons exercising control over the CAB.

### If the CAB is a legal entity that is wholly or partly owned by a larger organization, the CAB shall clearly document the activities, structure, and governance of that larger organization as well as its relationship with the CAB.

### If the CAB wholly or partly owns other legal entities, the CAB shall clearly define and document the activities and responsibilities of those other entities, as well as their legal and operational relationships with the CAB.

### In addition, a CAB that maintains multiple offices that perform some part of the regulatory review process shall ensure that the roles and responsibilities of the CAB and their other locations are defined and implemented.

### The CAB shall have legally enforceable arrangements with medical device manufacturers allowing personnel from the Regulatory Authority, which authorized their recognition, to assess the CAB’s activities associated with performing regulatory reviews. The agreement shall allow personnel from the Regulatory Authority access to records and documents pertaining to the manufacturer that are relevant to the regulatory review and decision-making process upon request.

### The CAB shall have legally enforceable arrangements with medical device manufacturers that will allow Regulatory Authorities to share all documents and records related to medical device regulatory reviews with other Regulatory Authorities, with which they have formal established confidentiality agreements covering provisions for protecting proprietary information and trade secret information.

### The agreements shall specify the responsibilities of the manufacturer and CAB, as well as the authority or responsibilities held by the recognizing Regulatory Authority, consistent with the laws and regulations in the relevant regulatory jurisdiction.

## Management of Impartiality

### The requirements of this section in no way preclude exchanges of technical or regulatory information between a CAB and a manufacturer.

**Specific Requirements for Medical Device CABs**

### The CAB shall be an entity that is independent of the manufacturer of the medical device for which it performs the regulatory review. The CAB shall also be independent of any other individual or group with an economic interest in the product as well as of any competitor of the manufacturer, such as distributors, authorized representatives, or importers.

### The CAB shall have documented and implemented a structure and procedures for safeguarding and promoting independence, objectivity, and impartiality throughout its organization, personnel, and regulatory reviews. These procedures shall effectively identify, investigate, and resolve any case in which a conflict of interests may arise. The CAB shall document any investigation, outcome, and resolution.

### The CAB shall ensure that its top-level management, the personnel responsible for carrying out the regulatory reviews, and the spouses (including domestic partners) and children of all these parties:

### are not involved in, or provide consultancy for, the design, manufacturing, assembly, marketing, supply, installation, distribution, import, purchase, or maintenance/servicing of the medical devices which they assess, nor act as an authorized representative of any of the parties engaged in those activities. They shall not engage in any activity that may conflict with their independence of judgment or integrity in relation to regulatory reviews, or undermine the confidence in their independence, impartiality, or objectivity; and

### do not use the services of any organization or individual that has provided consultancy services to the manufacturer, their authorized representative or a supplier being reviewed by the CAB, within a period of three years (or longer, if required by the recognizing Regulatory Authority) since the last consultancy services were rendered. Individual regulatory jurisdictions may have their own requirements regarding the impact of the timing of consultancy or other activities.

### Note: This does not preclude general training activities relating to medical device regulations, general marketing review, or related standards that are not manufacturer-specific.

### The CAB shall document any involvement in consultancy services in the field of medical devices undertaken by personnel prior to taking up employment with the CAB at the time of employment. The CAB shall monitor and resolve any potential conflicts of interest according to criteria set out in this document.

### If a CAB’s top-level management or the personnel responsible for performing the regulatory review were former employees of a medical device manufacturer or provided consultancy services in the field of medical devices for a specific manufacturer prior to taking up employment with the CAB, such personnel shall not be assigned to activities for that specific manufacturer or companies belonging to the same organization for a period of three years. Individual regulatory jurisdictions may have their own requirements regarding the impact of the timing of past consultancy or other activities.

### The CAB shall not advertise, commit to, guarantee, or imply the outcome of the regulatory review by the CAB on the basis of financial or other inducements.

### If the CAB is a legal entity that is wholly or partly owned by a larger organization, the requirements for impartiality in this document are applicable to both the CAB and the organization to which it belongs.

## Liability and Financing

**Specific Requirements for Medical Device CABs**

### The CAB shall take out professional liability insurance unless the government, in accordance with national law, covers liability.

### The justification for the scope and overall financial value of liability insurance shall be documented and include consideration of the level and geographic scope of activities of the CAB and the risk profile of the devices being assessed by the CAB.

## Non-Discriminatory Conditions

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Confidentiality

**Specific Requirements for Medical Device CABs**

### The CAB shall have documented procedures, and where applicable, equipment and facilities that ensure the secure handling of confidential information. The procedures shall ensure the confidentiality of information which comes into its possession during the regulatory review process.

### Except in relation to information which may be requested by a Regulatory Authority, the personnel of a CAB and any external regulatory reviewers, external technical experts, or outsourced organizations, shall not disclose information obtained in carrying out their tasks with respect to regulatory reviews with those who are not involved in the process.

## Publicly Available Information

### Exceptions to ISO/IEC 17065:2012

### For jurisdictions where the CAB issues the final marketing certification without the need for Regulatory Authority involvement, CABs shall make the marketing certification status of individual medical devices available upon request.

**Specific Requirements for Medical Device CABs**

### The information specified in ISO/IEC 17065:2012 Clause 4.6(a),(c), and (d) shall be made publicly available in all the geographical areas in which the CAB operates (e.g., web site), and not just available upon request.

### The CAB shall also maintain and make publicly available their regulatory review processes and their policy on impartiality, as well as the types of management systems in which it operates (see ISO/IEC 17065:2012 Clause 8).

### Where appropriate, the CAB must comply with specified Regulatory Authority requirements for the method of making information on the certified medical devices and manufacturer publicly accessible.

# Structural Requirements

## Organizational Structure and Top Management

**Specific Requirements for Medical Device CABs**

### The requirements listed in ISO/IEC 17065 Clause 5.1.3(d) and (e) also apply to implementation and oversight of any certification activities and requirements established by Regulatory Authorities.

### The CAB shall ensure that its personnel are current in practices and knowledge in relation to medical device technologies and regulatory requirements for conducting regulatory reviews of medical device marketing submissions.

### The CAB shall have the organizational capacity, including management, administrative support, and infrastructure, to undertake all contracted activities.

### The CAB shall actively participate in any regulatory coordination group activities established for the purpose of keeping the CAB’s personnel current regarding medical device legislation, guidance documents, standards, and best practice documents adopted in the applicable regulatory systems. The CAB shall identify a point of contact within their organization for each such activity.

### The CAB shall identify and apply relevant guidance and best practice documents, and justify why any documents that were identified and not applied were not considered relevant.

### The CAB shall establish a code of conduct for CABs in the field of medical devices (see Section 6.1.13 of this document). The code of conduct shall provide a mechanism for monitoring and verification of its implementation. Violations to the code of conduct must be investigated and appropriate action taken.

### The CAB shall have documented processes and procedures to ensure that personnel in the CAB do not make the final decision on their own evaluation. The recommendation and certification decision functions shall be performed by a person or persons independent of the evaluation team.

## Mechanism for Safeguarding Impartiality

**Specific Requirements for Medical Device CABs**

No specific requirements.

# Resource Requirements

## Certification Body Personnel

The resource and competence requirements for certification body personnel outlined in this section are general requirements. There may be more specific requirements for the regulatory authority having jurisdiction, including requirements related to:

* specific types of experts (e.g. clinical reviewers);
* whether certain types of personnel are internal or external resources;
* review; and
* the role of the reviewers and the final decision makers.

### *General*

**Specific Requirements for Medical Device CABs**

### A CAB shall comply with the specific requirements for the competence and maintenance of competence that can be found in IMDRF GRRP WG/N40 – “Competence, Training, and Conduct Requirements for Regulatory Reviewers,” and maintain current personnel records that include relevant competence, training, and conduct requirements as outlined therein.

### The CAB shall ensure that each regulatory reviewer, technical expert, or other personnel involved in the regulatory review process understands their duties, responsibilities, and authorities.

### The CAB shall have established processes for selecting, training, and formally authorizing regulatory reviewers and for selecting and familiarizing technical experts used in the certification activities. The initial competence evaluation of a regulatory reviewer shall include the ability to apply required knowledge and skills while performing regulatory reviews, as determined by a competent evaluator assessing the regulatory reviewer’s review.

### The CAB shall have a process to achieve and demonstrate effective regulatory reviews, including the use of regulatory reviewers possessing generic review skills and knowledge (see Section 7 of IMDRF/GRRP WG/N40), as well as technical experts possessing skills and knowledge appropriate for performing reviews in specific technical areas.

### The CAB shall ensure that regulatory reviewers and technical experts demonstrate competency regarding the CAB’s review processes, certification requirements and other relevant requirements, including regulatory requirements. The CAB shall give regulatory reviewers and technical experts access to an up-to-date set of documented procedures giving review instructions and all relevant information on the certification activities， including technical reference materials and relevant publications, regulations, standards, and guidelines.

### The CAB shall offer or provide access to specific training to ensure its regulatory reviewers, technical experts and other personnel involved in the regulatory review process are competent for the functions they perform. Additional guidance on training requirements are identified in IMDRF GRRP WG/N40.

### The final regulatory reviewer that makes the decision on granting, refusing, maintaining, renewing, suspending, restoring, or withdrawing certification, or on expanding or reducing the scope of certification, shall understand the applicable standard and certification requirements, and shall have demonstrated competence to evaluate the outcomes of the evaluation and recommendation of the review team.

### The personnel responsible for identifying competence requirements for the regulatory reviewer to perform specific regulatory reviews, and the final regulatory reviewer responsible for final review and decision-making on certification, shall be employees of the CAB and shall have proven knowledge and experience in the following:

1. the medical devices legislation, relevant guidance documents and standards adopted in the applicable regulatory system;
2. the types of qualifications, experience and expertise relevant to medical device regulatory reviews;
3. a broad base of medical device technologies, the medical device industry and the design and manufacture of medical devices;
4. the CAB’s quality management system, related procedures and the required qualification criteria; and
5. the training requirements for personnel involved in medical device regulatory reviews identified in IMDRF GRRP WG/N40.

***Management of Competence for Personnel Involved in the Certification Process***

**Specific Requirements for Medical Device CABs**

### A CAB shall have access to the necessary administrative, technical, and scientific personnel with technical knowledge and sufficient and appropriate experience relating to medical devices and the corresponding technologies.

### The management of the CAB shall have appropriate knowledge and processes to: set up and operate a system for the selection of the regulatory reviewers, technical experts, and other personnel involved in the regulatory review process; the verification of their competence; the assignment of their tasks; their initial and ongoing training; and, their instruction and monitoring to ensure that personnel who administer and perform the regulatory reviews are competent to fulfill the tasks required of them.

### The CAB shall identify at least one individual within its senior management having overall responsibility for all medical device regulatory reviews, within the scope of this document.

### A CAB shall be capable of carrying out all the tasks assigned to it with the highest degree of professional integrity and the requisite technical competence in the specific field, whether those tasks are carried out by the CAB itself or by other groups on the CAB’s behalf and under its responsibility.

### CABs are to ensure that regulatory reviewers, technical experts, and other personnel involved in regulatory reviews understand the importance of a code of conduct in maintaining integrity as discussed in IMDRF/GRRP WG/N40, Section 6.0. CABs are to keep signed statements of adherence to a code of conduct for personnel involved in regulatory reviews. The signed statement shall attest to at least the following elements included in the CAB’s code of conduct:

1. To act in a professional and ethical manner at all times.
2. To faithfully represent the interests of the recognizing Regulatory Authority(s) and CAB.
3. Not to act in any way prejudicial to the interests or reputation of the recognizing Regulatory Authority(s) and CAB.
4. Not to act in any way prejudicial to the integrity or objectives of the recognizing Regulatory Authority(s) and CAB.
5. To disclose any relationship, or financial interest, past or present, that may create a conflict of interest, or the appearance of a conflict of interest, and to notify management of any new conflicts of interest or potential conflicts of interest as soon as the case may arise.
6. Not to participate in any activity or relationship that may impair, or may appear to impair, one's objectivity, impartiality, or professional judgment.
7. Not to accept any inducement, gift, commission, discount or any other benefit not available to the general public from medical device manufacturers, their agents, their representatives, or economic operators.
8. To record and report truthfully and accurately regulatory review assessments in an impartial and unbiased way.
9. To record and report truthfully and accurately any material facts that may affect the reliability of the regulatory review process.
10. Not to provide any consulting services to manufacturers whose products are under regulatory review.
11. Not to disclose, verbally or written, any information obtained in the course of regulatory reviews to any third party, not including the recognizing Regulatory Authority(s), unless authorized in writing or required by law.
12. Not to use information obtained in the course of performing regulatory reviews for any personal gain.
13. Not to undertake any regulatory reviews for which one does not possess the required skills, knowledge or experience, formal designation or responsibility.
14. Not to undertake any regulatory reviews in a language where one is not proficient without the support of a translator and/or interpreter.
15. To continually improve one's proficiency, effectiveness, and quality of work.
16. To disclose to management, without delay, any breach of this statement by oneself or a colleague and to cooperate fully in the investigation of such a breach.

***Contract with Personnel***

**Specific Requirements for Medical Device CABs**

### CABs shall require regulatory reviewers, technical experts, and other personnel involved in the regulatory review process to sign a contract or document that declares any circumstances that could represent a potential conflict of interest, such as any association with the designer, manufacturer, supplier, installer, distributor, importer, purchaser, owner, or maintainer/servicer of the medical devices which they assess, or the authorized representative of any of those parties.

## Resources for Evaluation

***Internal Resources***

**Specific Requirements for Medical Device CABs**

### In addition to the specific standards listed in ISO/IEC 17065:2012 6.2.1 for testing, inspection, and management system auditing, the CAB shall meet additional applicable requirements such as regulatory, legal, and international standards; jurisdiction-specific and multilateral guidelines (including IMDRF documents); and the requirements outlined in this document.

***External Resources (Outsourcing)***

For the purposes of this clause, an external organization is one that is not subject to the CAB's quality management system.

**Specific Requirements for Medical Device CABs**

### In addition to the specific standards listed in ISO/IEC 17065:2012 6.2.1 for testing, inspection, and management system auditing, the bodies to whom the CAB outsources evaluation activities shall meet additional applicable requirements such as regulatory, legal, and international standards; jurisdiction-specific and multilateral guidelines (including IMDRF documents); and the requirements outlined in this document.

### Where a CAB uses an external organization for performing evaluation of marketing submissions, the CAB shall be responsible for identifying competence requirements for any regulatory reviewer, technical expert, or any other personnel involved in the evaluation process assigned to perform specific activities.

### Where a CAB uses an external organization for performing evaluation of marketing submissions, the CAB shall have adequate competence in such product areas to verify the appropriateness and validity of objective evidence provided by the external organization, to make the final review and decision on conformity to regulatory requirements.

### The CAB shall document all arrangements with external organizations involved in the regulatory review process and provide that information to the recognizing Regulatory Authority(s) on request. The CAB shall ensure that the external organizations allow the recognizing Regulatory Authority(s) to assess activities outsourced by the CAB.

### It is the CAB’s responsibility to ensure that all individuals utilized by an external organization that are involved in a regulatory review are directly assessed by the CAB to ensure consistency with IMDRF GRRP WG N40 and any assessment requirements.

### External resources shall not be responsible for the recommendation and certification decision.

# Process Requirements

## General

**Specific Requirements for Medical Device CABs**

### 7.1.1 The CAB shall establish procedures covering at least:

1. the manufacturer’s request for conducting a regulatory review;
2. the processing of such request, including the screening of the documentation, the qualification of the product as a medical device and its classification;
3. the acceptance or rejection of marketing submissions;
4. the language of the request, of the correspondence and of the documentation to be submitted;
5. where appropriate, the terms of the agreement with the manufacturer or authorized representative;
6. where appropriate, any fees to be charged for the regulatory review;
7. any requirement to perform a quality management system audit as part of the regulatory review;
8. the assignment of regulatory reviewers to a specific activity based on the regulatory reviewer’s competence to perform the regulatory review;
9. the decision-making process on the conformity of the marketing submission to applicable technical, legal, and regulatory requirements, including, where appropriate, the decision about issue, refusal, suspension, reinstatement, restriction, modification, or withdrawal of certificates as well as possible conditions or limitations to certificate validity;
10. the criteria for determining whether any change to the medical device is reportable to the Regulatory Authority having jurisdiction, and the assessment of reportable changes; and
11. where appropriate, the renewal of any certifications.

## Application

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Application Review (Screening)

**Specific Requirements for Medical Device CABs**

### The CAB shall screen the marketing submission to ensure that it contains the following:

1. information establishing the qualification of the product as a medical device;
2. the risk classification of the medical device;
3. the conformity assessment procedures applied by the manufacturer; and
4. sufficient information to demonstrate conformity with essential principles of safety and performance (per IMDRF GRRP WG/N47:2018 *Essential Principles of Safety and Performance of Medical Devices and IVD Medical Devices*), and any additional requirements specified by the recognizing Regulatory Authority(s).

The CAB shall also ensure that the information provided is relevant for the medical device under review.

### The CAB shall ensure that they have the necessary competence to review the marketing submission, including sufficient familiarity with all relevant regulations, standards, and guidelines related to the medical devices included in the marketing submission.

## Evaluation

**Exceptions to ISO/IEC 17065:2012**

In addition to the conditions described in ISO/IEC 17065:2012 Section 7.4.5 that state that the CAB can accept information related to certification completed prior to the submission for certification, the CAB can also accept information after the submission is received if agreed upon between the CAB and manufacturer.

**Specific Requirements for Medical Device CABs**

### The CAB shall evaluate the marketing submission according to the requirements specified by the recognizing Regulatory Authority(s). This evaluation can involve:

1. regulations, technical standards, and guidelines that are relevant to the medical device under review; and
2. relevantpublished IMDRF document**s,** including IMDRF GRRP WG/N47:2018 *Essential Principles of Safety and Performance of Medical Devices and IVD Medical Devices* andIMDRF GRRP WG/N52:2019 *Principles of Labelling for Medical Devices and IVD Medical Devices*, demonstrating conformity of the medical device under review with the relevant essential principles.

### The CAB shall ensure that the technical documentation provided in the marketing submission support the proposed classification of the medical device in the relevant regulatory jurisdiction.

### The CAB shall ensure that the technical documentation provided in the marketing submission support the proposed intended use of the medical device under review.

### If the CAB conducted an audit or receives the results of an audit as part of the marketing certification scheme, the CAB shall ensure that the audit results support the marketing submission.

## Review (Recommendation)

**Specific Requirements for Medical Device CABs**

### If the relevant Regulatory Authority requires QMS/GMP certification prior to marketing and an audit was not part of the CAB’s evaluation, the CAB shall ensure that the manufacturer holds appropriate QMS/GMP certification relevant to the medical device under review.

### The CAB shall document their recommendation for marketing certification in a marketing review report, based on the evaluation results. This report shall include information that addresses the requirements in section 7.7 of this document.

### If the CAB becomes aware of information included in the marketing submission that may require safety-related regulatory action (such as design changes that do not adequately address a postmarket issue involving marketed devices), such information shall be reported to the recognizing RA in writing within 5 working days.

## Certification Decision

**Exceptions to ISO/IEC 17065:2012**

For regulatory jurisdictions where the Regulatory Authority makes the final decision regarding the marketing of a medical device, ISO/IEC 17065:2012 Section 7.6 does not apply to the CAB. In these situations, the CAB shall submit the marketing review report discussed in Section 7.5.2 to the recognizing Regulatory Authority for the certification decision.

Beyond the requirements in ISO/IEC 17065:2012 Clause 7.6.3, the person(s) assigned by the CAB to make a certification decision shall be employed by the CAB.

**Specific Requirements for Medical Device CABs**

### A CAB conducting a regulatory review shall have sufficient and reliable evidence to support a decision of compliance or non-compliance with regulatory requirements.

## Certification Documentation

**Exceptions to ISO/IEC 17065:2012**

In regulatory jurisdictions where the final marketing decision must be made by the Regulatory Authority, the CAB shall provide their certification recommendation and marketing review report to the Regulatory Authority, and any certification documentation shall be issued by the Regulatory Authority in accordance with their legislation.

**Specific Requirements for Medical Device CABs**

### The CAB shall report marketing certifications to the recognizing Regulatory Authority when necessary, along with any other documentation required by the Regulatory Authority (which may include information regarding marketing submissions where marketing certification was not granted).

### When a CAB issues certificates and marketing review reports, they shall meet the requirements of the relevant Regulatory Authority.

### When a CAB issues reports and certificates intended for use by a specific Regulatory Authority, the reports and certificates shall accurately document:

1. the scope of the regulatory review;
2. the scope of certification issued, including clear identification of the certified medical device and its use; and
3. the review criteria used to assess the regulatory submission, including which Regulatory Authority requirements have been assessed.

## Directory of Certified Products

**Specific Requirements for Medical Device CABs**

### The CAB shall make its directory of certified products available to the recognizing Regulatory Authority.

## Surveillance

**Exceptions to ISO/IEC 17065:2012**

ISO/IEC 17065:2012 Section 7.9 does not apply.

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Changes Affecting Certification

**Specific Requirements for Medical Device CABs**

### If the regulatory requirements or guidelines change in regulatory jurisdiction where marketing certification was previously granted, the CAB shall revise their certification process to be consistent with these changes.

## Termination, Reduction, Suspension or Withdrawal of Certification

**Specific Requirements for Medical Device CABs**

### In addition to the requirements specified in ISO/IEC 17065:2012 Section 7.11, if the CAB is recommending termination, reduction, suspension, reinstatement, or withdrawal of certification without being requested by the manufacturer, they shall notify the recognizing Regulatory Authority within 5 working days. If required in the regulatory jurisdiction, final decisions regarding termination, reduction, suspension, reinstatement, or withdrawal of the certification shall be performed by the recognizing Regulatory Authority.

## Records

**Specific Requirements for Medical Device CABs**

### Records maintained by the CAB related to manufacturers shall include the following, in addition to the requirements in ISO/IEC 17065:2012:

1. marketing submission information and marketing review reports;
2. identification of any non-conformities identified during the certification process, and the actions taken to address these non-conformities;
3. committee deliberations and decisions, if applicable;
4. any correspondence with Regulatory Authorities regarding certification decisions or changes in certification; and
5. any certification documents, including the reports and certificates discussed in Section 7.7.2 of this document; and
6. any other relevant records necessary to establish the credibility of the certification.

### Records related to certification activities shall be retained for a time frame specified by the recognizing Regulatory Authority.

## Complaints and Appeals

**Specific Requirements for Medical Device CABs**

### The CAB shall forward to the recognizing Regulatory Authority(s) information on any complaint (e.g. whistleblowers) it receives about a medical device manufacturer that could indicate an issue related to the safety and performance of medical devices or a public health risk.

### Any appeals regarding a regulatory review performed by the CAB shall be handled by the CAB. The CAB shall notify the recognizing Regulatory Authority of any decision made regarding these appeals that changes the final review decision. The recognizing Regulatory Authorities may also establish their own process for further appeals, subsequent to appeals made to the CAB.

#  Management System Requirements for Medical Device CABs

## Options

**Specific Requirements for Medical Device CABs**

### The CAB shall establish, document, implement, maintain, and operate a quality management system that is appropriate to the nature, area, and scale of its reviews and capable of supporting and demonstrating the consistent achievement of the requirements of applicable medical device legislation or regulatory policies/programs.

### The CAB shall retain records of conformity to this document for a period of time not less than 15 years from the creation of the record.

### The CAB shall perform measuring, monitoring and the analysis of their review program to provide information relating to the characteristics and trends of their processes such as: consistency in marketing review reports, feedback from medical device manufacturers, etc.

### The CAB, when adhering to Option A or Option B under Clause 8 of ISO/IEC 17065:2012, shall conduct internal audits that cover its organization’s structure and activities at all locations involved in medical device regulatory reviews.

## General Management System Requirements (Option A)

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Control of Documents (Option A)

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Control of Records (Option A)

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Management Review (Option A)

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Internal Audits (Option A)

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Corrective Actions (Option A)

**Specific Requirements for Medical Device CABs**

No specific requirements.

## Preventive Actions (Option A)

**Specific Requirements for Medical Device CABs**

No specific requirements.

# Information Requirements

## Information Exchange Between the CAB and Recognizing Regulatory Authority(s)

### The CAB shall designate a function within its organization for timely exchange of information with recognizing Regulatory Authority(s) or to respond to enquiries from recognizing Regulatory Authority(s). The CAB shall fulfill this function at all times.

### The CAB shall report to the recognizing Regulatory Authority(s) that performed its initial recognition when it becomes aware of any fraudulent activities or counterfeit products. The CAB is not responsible for establishing objective evidence but must report such activities or products in writing within 5 working days from the date of discovery.

### The CAB shall provide information to the recognizing Regulatory Authority(s) about any regulatory reviews and decisions to grant or refuse marketing certification. Recognizing Regulatory Authorities may have their own requirements regarding the timing and mechanism for providing this information.

### The CAB shall notify the recognizing Regulatory Authority(s) in writing within 5 working days from the date of a decision to terminate, reduce, suspend, reinstate, or withdraw marketing certification. The notification shall include a rationale for such action.

### The CAB shall notify the recognizing Regulatory Authority(s) in writing within 5 working days of changes in matters that may affect the abilities of the CAB to fulfill the requirements for recognition, such as:

1. its legal, commercial, organizational or ownership status;
2. the points of contact for regulatory coordination group activities discussed in Section 5.1.4 of this document;
3. address(es) of the CAB; or
4. scope of recognition as a CAB.