# REVISION OF THE CONTINUOUS IMPROVEMENT PROCEDURE (CIP) ESC-PRO-30-011 V1-2

Webinar 16 January 2025

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# **PURPOSE OF THE WEBINAR**



### 1. Informative

Help understand the changes and the rationale behind them

### 2. Question and feedback

Encourage clarification related questions.

For feedback, please use the consultation platform.

For general enquiries <a href="mailto:systemdevelopment@fsc.org">systemdevelopment@fsc.org</a>







- Why we are revising the CIP.
- **2.** Editorial changes to the CIP.
- **3.** Changes to closure of nonconformities
- **4.** Future structure of FSCs CB standards.
- **5.** CIP revision timeline.



### **1. Why are we revising the CIP procedure?**



Need to align with ISO requirements and the German National Accreditation Body "DAkkS" Several nonconformities for FSC were issued, to be addressed in the accreditation standards. Core issues are the following:

Closure of nonconformities	Evaluations are done in a consistent manner	Certification requirements are separated from accreditation requirements	777
Addressed in FSC- STD-20-001 Also issue with Continuous Improvement Procedure.	Addressed in all FSC accreditation standards.	Needs to be implemented for: 1. Continuous Improvement Procedure 2. Ecosystem Services Procedure 3. Regulatory Module	

# 2. Editorial changes



The Organization instead of "CIP user".	Other changes in terminology.	Editorial changes in the Introduction.
Reference to "The Organization" instead of "CIP user" to align with the normative framework.	Change in terminology, e.g. Criteria designated as low risk "are not applicable" in case of inactive management units instead of stating "conformity is not required to be demonstrated".	Editorial changes in the Introduction to highlight the FSC requirements for FM certification for SLIMF and CF applying the CIP: Core Criteria and Action Plan



### 3. Closure of Nonconformities (1/3) Relevant requirements from ISO 17065





#### **Nonconformities**

ISO does not distinguish between minor and major and only focuses on corrections and not on the prevention of reoccurrence (corrective action).



### For granting certification

The certification body shall only grant certification when:

- Decision to grant has been made;
- the client has fulfilled the certification
  requirements (Source 7.7.3).



#### For surveillance

The surveillance shall include the certification decision (Source 7.7.3).

When a nonconformity is identified, CB shall consider and decide upon appropriate action (increased surveillance, scope reduction, suspension, withdrawal) (Source 7.11.1).

# 3. Closure of Nonconformities (2/3) Latest draft FSC-STD-20-001 requirements



#### Definitions

- **Correction:** action to eliminate a nonconformity
- **Corrective action:** action to eliminate the cause of a nonconformity and to prevent recurrence.

#### **Main evaluation**

### Minor and Major nonconformities:

 Close all nonconformities: corrections and corrective actions before certification is granted



## 3. Closure of Nonconformities (3/3) Latest draft FSC-STD-20-001 requirements

#### Surveillance

### Major non-conformities

- Corrections
- Corrective actions
- Timeline: CBs get 1 month to develop the nonconformity report
- CHs: within three months from receiving the nonconformity report

#### Surveillance

### **Minor non-conformities**

• Corrections

Same timelines as for major nonconformities

Corrective actions

Maximum timeline is until the next surveillance evaluation

Current CIP version allows minor NCs to be corrected within 2 years.

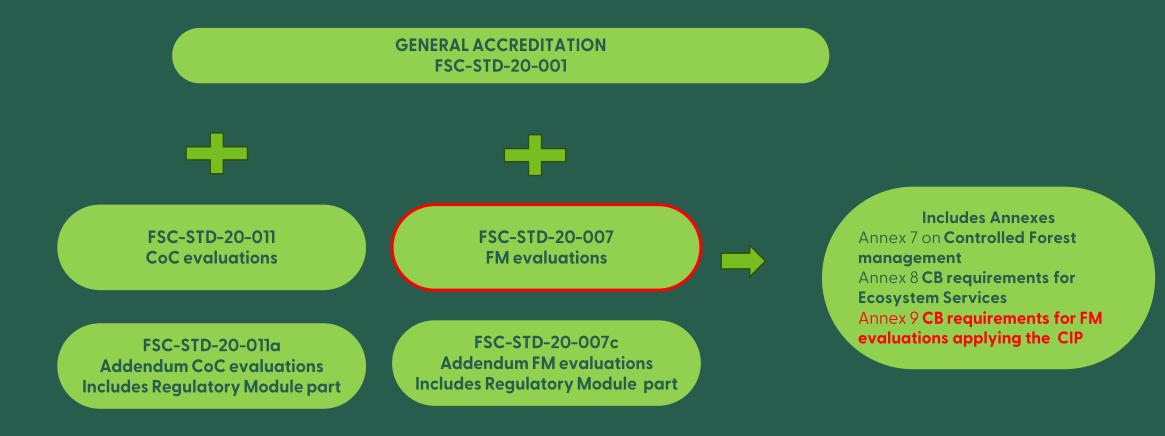




# 4. Future structure of FSCs CB standards

Certification requirements are separated from accreditation requirements





### **5. CIP revision timeline**

Category	Activity
Process type	A joint accelerated process is applied, resulting in FSC-STD-20-007 V4-1, and FSC-PRO-30-011 V1-2
Objective of revision process	To align with FSC-STD-20-001 and to address DAkkS nonconformities
Timeline	Aligned with FSC-STD-20-001 revision process





# Thank you



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**FSC®** International Center

Adenauerallee 134, 53113 Bonn, Germany T +49 (0) 228 367 66-0 F +49 (0) 228 367 66-65 FSC International Center © All rights reserved FSC® F000100

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