

CELEBRATING 70 YEARS OF THE CHICAGO CONVENTION

# Workshop on quality assurance for the implementation of an instrument flight procedure

29 March -2 April 2021



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# Flight Procedure Design Regulatory Approval (Doc. 9906, vol. 1)





- 1. Regulatory framework
- 2. State responsibilities
- 3. Flight procedure implementation diagram
- 4. Flight procedure regulatory approval (requirements & approval)
- 5. Best practices



### **Regulatory framework**



- There is no dedicated ICAO Document on IFP regulatory approval
- Provisions provided in various documents

### Regulatory framework

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### State responsibilities

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- □ The implementation of flight procedures is the responsibility of States!!!
  - © Establish operating rules for IFPD;
  - <sup>C</sup> Carry out FPD process or delegation to third parties.
- Establish a quality system for the entire IFP process:
  - <sup>CP</sup> Ensure their quality and safety through:
    - Review, verification, coordination, validation of certain points, etc.

Approve the instrument flight procedures prior to validation.

### State responsibilities

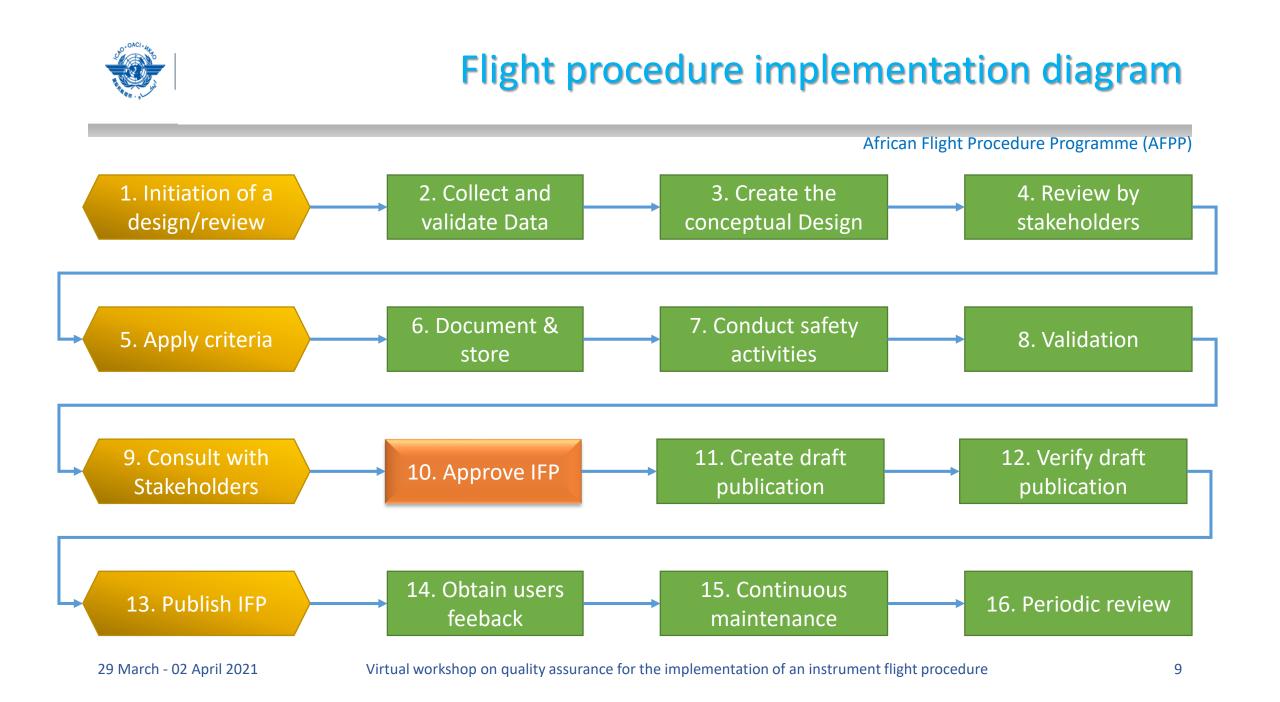


Initiation step (1)	<ul> <li>Each State should describe the initiation and submission process valid within their State;</li> <li>Establish interval for periodic review (5 years maximum).</li> </ul>
Apply criteria (5)	• Establish applicable design criteria
Document and store (6)	• Define minimum period of time during which the withdrawn IFP documentation should be kept.
Validation	• Establish measures for the validation of IFPs
Approve the IFP (10)	• Approve IFP prior to publication.
Publication	<ul> <li>Publication of the IFP</li> <li>Delegation possible</li> </ul>
Periodic review (16)	• Establish interval for periodic review (5 years maximum).

### **State responsibilities**



IFP designers	<ul> <li>Each State should standards for the required competency level for the IFP designers;</li> <li>Ensure that designers have acquired and maintain this competency.</li> </ul>
Training	<ul> <li>Approve training courses/programme conducted by Procedure Design Service Providers (PDSP);</li> <li>Establish minimum qualifications and training for Flight Validation Pilots</li> </ul>
Flight Validation Pilots	<ul> <li>Each State should standards for the required competency level for the FVPs;</li> <li>Ensure that FVPs have acquired and maintain this competency.</li> </ul>



#### 10

### Flight procedure regulatory approval

- There is no specific guidance for the IFP regulatory approval:
   Each State to develop its own.
- However, approval process may consider many aspects:
   The actors qualifications;
  - The software tools used;
  - The completeness of the flight procedure approval request bundle;
  - The compliance with the respective quality assurance steps;
  - The compliance with the data requirements, the design criteria and the annex 4 charting criteria.







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#### The actors qualifications: the Designers

□ Many questions:

- Is there any required license for the designers?
- Is the designer adequately trained and is he maintaining his competency?
- What is his experience vs the complexity of the IFPs to be designed?





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#### The actors qualifications: the FVPs

#### Ensure:

- The license of the FVP;
- His level of qualification and experience.





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#### The software used

Ensure:

- That the software tools used are validated:
  - Flight Procedure Design tools
  - Charting tools;
  - Ground validation tool;
  - Simulator evaluation and/or flight evaluation tools;
  - Etc.



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# Flight procedure regulatory approval

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#### **Completeness of the approval request package**



- The quality record documents:
  - Approved conceptual design (step 4);
  - Results of the validation (step 8);
  - Stakeholders endorsement (step 9).
- Some of the most important outputs (from the steps).





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#### **Example of regulatory approval package**

#	Document	Provenance	Step
1	Managerial decision to set up the design	Output	Step 1
2	Data collection report. Content: • Stakeholders requirements • Summary of data collected		Step 2
3	<ul> <li>Stakeholders review report:</li> <li>Formally approved Conceptual Design (CD) or</li> <li>Decision to discontinue;</li> <li>Planned AIRAC implementation date (resources, training constraints, etc.)</li> </ul>	Quality record	Step 4



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#### **Example of regulatory approval package**

#	Document	Provenance	Step
4	<ul> <li>Technical report containing:</li> <li>All strategies, calculations, forms, etc.;</li> <li>Draft charts and draft coding tables (if applicable)</li> <li>List of waypoints</li> <li>Etc.</li> </ul>	Output	Step 6
5	<ul> <li>Safety assessment report. Content:</li> <li>Amplitude of the study case;</li> <li>Identified dangers;</li> <li>Mitigation measures, etc.</li> </ul>	Output	Step 7



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#### **Example of regulatory approval package**

#	Document	Requirement	Step
6	<ul> <li>Validation report:</li> <li>Ground validation report;</li> <li>Simulator evaluation report (if applicable)</li> <li>Flight evaluation report (if applicable)</li> </ul>	Output Quality record	Step 8
7	<ul><li>Stakeholder consultation report. Content:</li><li>Stakeholders endorsement.</li></ul>	Output Quality record	Step 9





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#### Best practices can consist on (example):

- Establishing clear PANS-OPS regulation:
  - Data requirements, Design criteria, QMS, Charting, Validation, etc.
- Setablishing a PANS-OPS implementation team on the CAA side for each design project:
  - The PANS-OPS team follow the project from initiation;
  - The team :
    - Checks the actors qualifications and competencies (Designers, FVPs);
    - Ensure the validity of the software used (Design, charting, etc.);
    - Reviews the approval request bundle to ensure:
      - ✓ Completeness of the request bundles;
      - ✓ Adherence to the quality assurance process;
      - $\checkmark$  Compliance with the applicable regulation.



**Best practices** 

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#### **Best practices can consist on (example):**

#### The inspectors may:

- Develop approval procedures and processes;
- Job aids.
- Example of job aid.



