# Public Review Draft 1 of the Biometric iTC PP-Module & Supporting Document v1.1 Overview

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## Introduction

This is an announcement of a public review period for the PP-Module and PP-Module Supporting Document from the BIO-iTC. By design these documents do not provide information about the methodology for performing Presentation Attack Detection. These are being handled as a toolkit which can be updated separately (and ideally more rapidly) without needing changes to the PP-Module and Supporting Document.

This document will provide information about where to find the documents, how to provide feedback and information about the current status of the documents.

Please email isec-itc-bio-info@ipa.go.jp if there are any questions about the Review process.

The latest status of the Review Period, including the latest copies of any documents, can be found at the Biometrics Security iTC page.

## **Review Status Phase**

The documents listed here are in the v1.1 **Public Review Draft 1** phase.

#### **Publication Date**

July 1, 2021

#### **End of Comment Period**

August 15, 2021

## **Documents for Review**

The following are the documents included in this v1.1 Public Review Draft 1 Period:

Table 1. Review Documents

Title	Version	Link
collaborative PP-Module for Mobile biometric enrolment and verification - for unlocking the device -	1.1	Download PP-Module
PP-Configuration for Protection Profile for Mobile Device Fundamentals and collaborative Protection Profile Module for Mobile biometric enrolment and verification - for unlocking the device -	1.1	Download PP-Configuration
Supporting Document Mandatory Technical Document: Evaluation Activities for collaborative PP-Module for Mobile biometric enrolment and verification - for unlocking the device -	1.1	Download Supporting Document

The following are the documents to use in support of the Public Review Draft 1 Period.

Table 2. Supporting Documents

Title	Link
Comment Matrix	Download Comment Matrix

# **Document Release Process**

The BIO-iTC follows the document release process below for the publication of documents. Generally the PP (PP-Module and PP-Configuration) and SD would be released at the same time.

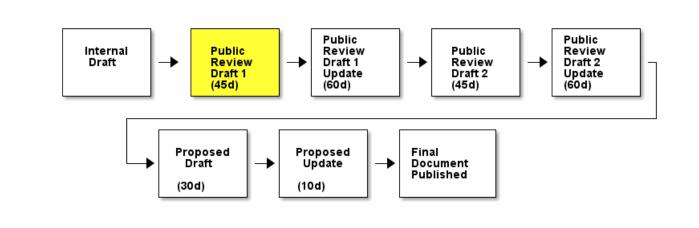


Table 3. Review Timeline

Phase	Time	Description
Internal Draft		The normal, pre-release process for creating the documents
Public Review Draft 1	45 days	iTC has voted according to Terms of Reference to release this version for public review. Public (i.e. from non-iTC participants) comments are accepted during this period
Public Review Draft 1 Update	Up to 60 days	The iTC will review all received comments and update the documents accordingly
Public Review Draft 2	45 days	iTC has voted according to Terms of Reference to release this version for public review. Public (i.e. from non-iTC participants) comments are accepted during this period  PAD Toolbox drafts will be published with this review (final review on toolboxes may proceed independently from rest of documents).
Public Review Draft 2 Update	Up to 60 days	The iTC will review all received comments and update the documents accordingly
Public Review Draft 3 (Optional, not shown)	45 days	iTC has voted according to Terms of Reference to release this version for public review. Public (i.e. from non-iTC participants) comments are accepted during this period
Public Review Draft 3 Update (Optional, not shown)	Up to 60 days	The iTC will review all received comments and update the documents accordingly
Proposed Draft	30 days	iTC has voted according to Terms of Reference to propose this as the final document. Public (i.e. from non-iTC participants) comments are accepted during this period

Phase	Time	Description
Proposed Update	10 days	iTC reviews any further comments and prepares the document for final publishing (updating all dates, producing official versions for publication)
Final Document Published		Documents are posted to Common Criteria Portal

The iTC may decide, based on the comments received during the Public Review Draft 1 period, that a Public Review Draft 2 period is needed. Public announcement of a second review draft or a proposed draft will be made once all comments have been addressed.

## **Review Process**

There are two ways to contribute comments and suggestions to the iTC. The first is through GitHub, the second by spreadsheet. It should be noted however that comments that are received via the spreadsheet will be added to the GitHub platform to allow for a comprehensive discussion. Also, feedback for comments is only provided via the answers in the GitHub comments.

Each comment should have a suggested resolution be proposed if a change is needed to the document.

### **GitHub Review Process**

To use GitHub to submit comments, you must have a GitHub account (and it is assumed you know how to use GitHub). Each comment should be submitted as an individual Issue with the Label "Public Review" assigned. Pull Requests created for any issues will be linked to these Issues for traceability.

## **Comment Matrix Spreadsheet Review Process**

In the Table 2, "Supporting Documents" table there is a link to the Comment Matrix spreadsheet. There are instructions for using the Matrix on the second worksheet. Please create a separate copy of the spreadsheet for each document.

Email the spreadsheets to isec-itc-bio-info@ipa.go.jp.

## **Additional Notes**

Among all the changes made to the documents, there are three particular areas where the BIO-iTC is asking for specific review and comments. These areas are noted below.

## Addition of IAPAR for FIA\_MBV\_EXT.3

The FIA\_MBV\_EXT.3 requirement has been updated to include Imposter Attack Presentation Accept Rate (IAPAR) to enhance the specific requirements around presentation attack. There are several

questions about this addition the BIO-iTC is looking for feedback on.

- Is the maximum value of 15% (the worst failure rate) an acceptable limit?
- Is there a different rate that would be preferred (one concern is that the requirements are for a whole system and not components, so any measure should be targeted to the whole system)?
- Is there a similar rate that would be preferred for use with enrolment? IAPAR is not valid for enrolment, and the BIO-iTC was unaware of any other measure for the enrolment process.

## FPT\_BDP\_EXT.1 Evaluation Activities for ATE\_IND.1

The PP-Module has been updated to better integrate with the PP\_MD (Mobile Device Fundamenmtals), and the FPT\_BDP\_EXT.\* requirements have all be adjusted. The remaining requirement is targeted to ensuring that biometric data is not sent through the main OS on the device. The current ATE\_IND.1 testing strategy is written with the FCS\_CKM.1 test activities in mind, where memory is dumped during the biometric capture process to check that the capture isn't accessible to the main OS itself.

The BIO-iTC is looking for comments about the best method for performing this type of test, or even if this should be proposed as a test strategy. An alternative approach that has been discussed is leaving this as a documentation review where the vendor would provide an architecture description of how the biometric system interacts with the rest of the device (including the main OS).

# **Biometric Sample Quality Assessment**

The FIA\_MBV\_EXT.2 and FIA\_MBE\_EXT.2 requirements have been updated to point to a specific quality metric, either public (such as ISO 29794-1) or one defined by the developer. While the expectations for meeting a public assessment seem pretty clear (that the report of that review be provided), there are questions about what should be used as the requirements for a developer defined quality assessment.

A concern has been raised that mobile devices (the first target for the BIOPPM) are not normally tested/reviewed against the public quality measures and usually focus on a balance that meets the FAR/FRR needs of the device, and that the public measures are not designed for mobile devices in general, but other systems that may share biometric data.

The BIOSD has a section with a proposed outline of what needs to be provided to the evaluator for review, and is looking for comments on this section to better target it to both what a validator would expect to see as complete and also what a vendor would consider appropriate given the nature of the quality measures.

It is important to note that this information is needed for testing that the samples provided to the biometric system have sufficient information needed to build templates which will result in the desired performance, i.e. acceptable error rates. So the focus is on how to create poor samples and how to ensure they are discarded from use.