# **IV&V** Quality Manual

Version: S

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Note: The official version of this document is maintained in IV&V's internal IV&V Management System Website (<a href="https://confluence.ivv.nasa.gov:8445/display/IMS">https://confluence.ivv.nasa.gov:8445/display/IMS</a>). This document is uncontrolled when printed.

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## **Purpose**

This document defines the manner in which the NASA IV&V Program implements the NASA IV&V Management System (IMS) as a quality management system.

The IMS is designed to meet the requirements of the International Organization for Standardization (ISO) 9001: 2015 Standard, the ISO 17020:2012 Standard, and any other requirements for a Federal Risk and Authorization Management Program (FedRAMP) Third Party Assessment Organization (3PAO).

## Scope

This document applies to the work performed under the scope of the NASA IV&V Program's ISO certification:

- Independent Software Verification and Validation
- System Software Assessments
- Systems and Software Engineering Research
- Software Support for the Office of Safety and Mission Assurance (OSMA)

This scope encompasses all of the work activities performed by the NASA IV&V Program and documented in the IMS. The IMS represents a foundational component of the IV&V Program that enables the NASA IV&V Program's mission and efforts to achieve its vision.

Understanding the key elements of the NASA IV&V Program's mission helps employees to advance that mission across several primary areas of focus, known as functional organizations.

The functional organizations and their functional statements can be found here: IV&V Program Implementation and Reporting

## **IV&V Management System**

#### General

The IMS encompasses activities that affect the products and services that the NASA IV&V Program provides to customers. The IMS contains processes, procedures, work instructions and supporting documentation that *ensure high quality products and services are provided by the NASA IV&V Program. The IMS enables and standardizes* the planning, performance, control, and measurement of NASA IV&V operations. The NASA IV&V Program practices continuous improvement in its refinement and enhancement of the IMS.

The IMS is documented and structured in the following manner:

#### **IMS Documentation**

The following figure the IMS structure.

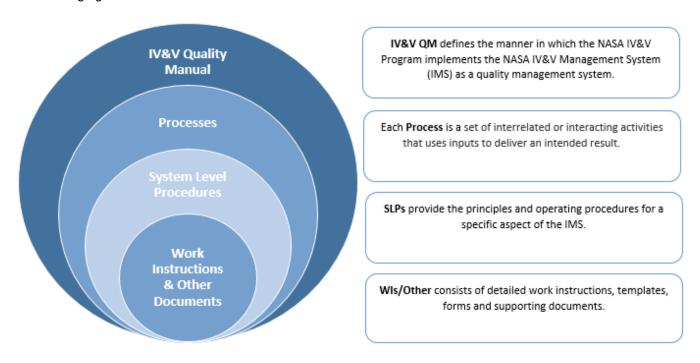


Figure 2 - IVV QM - IMS Documentation Structure.docx

Figure 2 – IMS Documentation Structure

The IMS Documentation Master List in the IMS Confluence pages provides a current representation of all content within the IMS including process/document owners and revision information.

### **Quality Policy**

The NASA IV&V Program's Vision statement and Mission statement serve as the Program's quality policy. These statements and the work performed by the IV&V Program are driven by agency-level content.

The NASA IV&V Program's Vision and Mission statements and key agency-level drivers can be found here: https://confluence.ivv.nasa.gov:8445/pages/viewpage.action?pageId=123602129

Additional information regarding how the IV&V Program meets ISO requirements can be found here: https://ecm.msw.faircon.net/OTCS/llisapi.dll/open/6953205

### **Quality Objectives**

The NASA Strategic Plan outlines the strategic direction, goals, and priorities NASA IV&V will pursue to achieve NASA's Vision and Mission. The IV&V Program is a NASA Program within the Office of Safety and Mission Assurance and established in accordance with NPD 1000.3. The Program's quality objectives consist of the NASA IV&V's overall responsibilities that are outlined in NPD 1000.3E.

The NASA Strategic Plan can be found here: https://confluence.ivv.nasa.gov:8445/pages/viewpage.action?pageId=141101084

The NASA IV&V Quality Objectives/NPD 1000.3E can be found here: https://confluence.ivv.nasa.gov:8445/pages/viewpage.action?pageId=141101068

# **Appendices**

The following appendices contain content maintained in separate documents. Follow the links provided to view the documents.

Appendix A – Acronyms and Definitions

Appendix B – Responsibilities and Authorities

## **Version History**

VERSION HISTORY				
V e r s i	Description of Change  Rationale for Change	A u t h or	E ff e c ti v e D a te	

B a sic	Initial release to replace Ames QM	S ia m a k Y a s si ni I T / 3 32	0 9 / 0 9 / 1 9 99
A - H	Older revision information may be located in the Version History Overflow Document	V a ri o us	0 5 / 0 7 / 2 0 0 1 - 0 1 / 2 4 / 2 0 066
1	Updated content; moved appendix material to separate documents	S t e p h a ni e F e r g u s on	1 0 / 1 0 / 2 0 07

J	Updated Vision, Quality Policy, and document references; updated terms and definitions resulting from re-engineering process	S t e p h a ni e F e r g u s on	0 4 / 0 8 / 2 0 08
К	Updated terminology to reflect the current organizational structure and to meet the ISO 9001:2008 Standard	S t e p h a ni e F e r g u s on	0 7 / 1 4 / 2 0 09
L	Updated to reflect new organizational structure	S t e p h a ni e F e r g u s on	0 7 / 1 3 / 2 0 10

M	Revised information on PEP; added execution plan process diagram		G r e g B la n ey	0 8 / 3 1 / 2 0 10
N	Updated the Quality Policy, Leadership direction and Quality Objectives. Sections 1.2, 1.2.1, 1.2.2, and 1.3 are affected.  Added IVV 09-9 to section 4.3.  Added IVV 03 and IVV 26 to section 4.5.		G r e g B la n ey	1 1 / 2 4 / 2 0 10
O	Updated Knowledge Management and KM roles, Public Affairs, etc.	Updated to accurately reflect updated roles and organizational changes	N a t al ie A lv a ro	1 0 / 2 4 / 2 0 11
P	Updated vision/mission, add description of Strategic Plan, remove details regarding office ownership of SLP's, remove detailed office descriptions, add description of Office Execution Planning process (and the tie between that content and the Strategic Plan, including Outcomes serving as quality objectives), removal of GSFC and HQ org charts.	Sync the QM with the IV&V Program Strategic Plan (approved Sept. 2012) (addresses PAR # 2012-P-376), simplify the QM by removing content that isn't required and describes our organization in general (rather than describing our organization's implementation of a quality management system).	J e ff r e y N o rt h ey	0 3 / 2 7 / 2 0 13

Q	Modified Section 3.2 to clarify that Section 7.6 (of the ISO 9001:2008 Standard), Control of monitoring and measuring equipment, is excluded.	Equipment is not used by the IV&V Program to provide evidence of conformity of product to determined requirements. (PAR: 2014-P-396).	J e ff r e y N o rt h ey	0 8 / 2 8 / 2 0 14
R	Added language to Purpose to be explicit about our intent. We want a single management system that meets all requirements. Under IMS Documentation - updated the terms to re-use exiting content and removed content not needed to be called out certain required sections for the standard.	To address outdated references to the ISO 9001:2008 standard. (PAR #2017-P-498)	J e ff N o rt h	1 1 / 2 8 / 2 0 18
S	Removed references to the IV&V Program Strategic Plan, added link to functional organizations and their functional statements, removed organizational structure chart, removed management responsibility, vision statement, and mission statement, updated Quality Policy to be the Program's Vision and Mission statements and added a link to the statements, updated the Quality Objectives to be the responsibilities outlined in NPD 1000.3E and added a link to the NPD, added a link to the Agency's Strategic Plan, removed reference to Office Execution Plans (OEP), removed the NASA IV&V Execution Plan Model and description, removed organization description, responsibility and authority, and management review description of Quarterly Metric Reviews (QMR), removed figure for process for customer feedback.	This Program-level content is already being maintained and is available to our program in Confluence. The compliance matrix will now capture the information on how IV&V meets the ISO requirements.	A le x A y e rs	0 8 / 0 5 / 2 0 20