

QUALITY MANUAL

Effectivity Date

Revision Level 2

Document Code TIEZA.QM.03

DOCUMENT MANAGEMENT

3. TIEZA's QMS Documents

3.1 General

TIEZA's QMS documentation include the quality manual, the documented procedures required by the ISO 9001 standard, and internal and external maintained documented information determined by TIEZA to be necessary for the effective QMS planning and operation.

This maintained documented information also contains appropriate criteria for determining whether prescribed activities have been completed satisfactorily.

The documentation is presented in the following hierarchy:

| Level | Type of Document | Description |
|-------|----------------------------|--|
| 1 | Quality Manual (QM) | The highest level of QMS documents is the Quality Manual (QM). The QM describes, in the broad terms, the overall adaptation of ISO 9001 in TIEZA. The QM describes how each applicable section of the standard is to be implemented to meet the requirements of certification bodies and customers. |
| | | The QM contains the scope of TIEZA's quality management system, including details of and justification for any exclusion in the ISO 9001 standard. This QM describes the sequence and the interaction of the QMS processes. This QM also establishes the structure, authority, and responsibility for the maintenance of the QMS. |
| 2 | Quality Procedures (QP) | The Quality Procedures (QP) are documents stating what and how things should be done, the materials and/or equipment that should be done, used, and who will do it. The QP address applicable requirements and QMS policies which are reviewed and approved by authorized personnel. The QP contain the purpose for the policy. It must contain the scope of the policy and the definition |



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| Level | Type of Document | ent Description | |
|-------|---------------------------|---|--|
| | | of terms. These are established to support the processes described in the QM. | |
| 3 | Work Instructions (WI) | Work Instructions (WI) provide detailed steps to conduct specific work activities of each offices and are prepared as needed to supplement procedure requirements and to ensure that critical work scopes are carried out in a consistent manner. This includes office standard operating procedures. | |
| 4 | Quality Records | A. External These are documented information generated from outside TIEZA and used as reference to ensure effective planning and control of its QMS These may include Department Orders Memorandum Circulars, technical manuals, etc. | |

3.2 Control of Documents

The authority has established a documented procedure which ensures the necessary control on the approval, issuance, review, update and reapproval of documented information, and data affecting the processes within the scope of the QMS.

B. Internal

requirements.

- a. Creation/ Revision of Documented Information
- b. Pre-registration of the Documented Information
- c. Review and Approval of the Documented Information
- d. Registration of the Documented Information
- e. Distribution and Maintenance of Documented Information
- f. Online Document Management System QMS Module

Refer to TIEZA.QP.01 (Control of Documents)



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DISCLAIMER:

These are documented information that furnish evidence of conformity of products, services, and/or activities rendered to TIEZA's QMS

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Documented Information and Data Approval and Issue

During document preparation and review, the authorized personnel, reviews the documented information to ensure that the requirements can be met within a timely manner once the documented information is formally issued.

All levels of management are responsible for assigning responsibilities to ensure that documented information and data are controlled in accordance with established procedures and resolving issues pertaining to policy and procedure content, application, and use.

Moreover, due to the nature of the organization, there are several external documented information that TIEZA must use and follow, but which must be controlled. Access to and use of external documented information is in accordance with applicable TIEZA policies.

Document and Data Changes

The QMS contains a documented procedure to ensure that procedures are identified, prepared, documented, and revised in a uniform manner, and that they give clear guidance and direction to their users.

Change to procedures, instructions and drawings are approved and documented prior to implementation, and are made available at the location where the activity will be performed prior to commencing work.

Invalid or obsolete documented information are promptly removed from all points of issue or use, or otherwise assured against unintended use. Any obsolete documented information that are preserved are identified and archived.

3.3 Control of Record

The Authority has established and maintained a documented procedure which defines the controls of documented information. This includes

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requirements and responsibilities needed for the following (electronic and/or hard copies):

- Classification/ Identification
- Legibility
- Collection
- Indexing
- Filing
- Access
- Storage
- Protection
- Distribution
- Retention
- Maintenance
- Retrieval and
- Disposition of all records

The documented information established by each office are retained to provide evidence of conformity to requirements and of the effective operation of the QMS.

All documented information that are retained are kept and secured in a safe place protected from damage, loss, and deterioration.

All documented information beyond the retention period shall be disposed accordingly based on National Archives of the Philippines (NAP) guidelines.

- a. TIEZA Inventory of Records
- b. Agency Records Disposition Schedule

Refer to TIEZA.QP.03 (Control of Records) and Republic Act 9470: Implementing Rules and Regulations of RA 9470, otherwise known as the National Archives of the Philippines Act of 2007.

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3.4 Document Formatting

3.4.1 Standard content

| Procedure | | Work Instruction | |
|----------------------------|--|---|--|
| a. b. c. d. e. f. g. h. i. | Scope Definition of Terms (alphabetized, terms-bold) Acronym Guidelines Procedure Flow Procedure Details | a. Purpose b. Scope c. Guidelines d. Process Flow e. Process Details f. References g. Form/s (if necessary) | |

- 3.4.2 The **header and footer** used for this document is the official template to be used for the quality manual, quality procedures, and work instructions.
- 3.4.3 The **header** bears the **official logo of TIEZA** with a size of 1.25" by 1.25", located at the upper left corner. The official template will be provided only by the Document Controller (DC) in doc format.

The **Header** consists of the following:

- a) TIEZA Logo
- b) TIEZA name
- c) Title
- d) Level/Kind of Document
- e) **Effectivity date
- f) **Revision level -starting with 0 for the creation
- g) Document Code (see #3.4.8)

** to be accomplished by Document Controller

Date: 0/6/mv
DC Signature: Pd
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3.4.4 The **footer** consists of portion for the stamp of the DC for Master copy and Controlled copy, the disclaimer and the **page number**: (example: Page 1 of 13).

3.4.5 Official font and size of contents is Calibri 14.

3.4.6 Official paper size is A4 -8.27" x 11.69".

3.4.7 Margin: Top-0.5", Bottom/Left/Right-1.0", Header-0.5", Footer-0.5"

3.4.8 Document Coding:

| TIEZA WIDE | | SECTORAL/ DEPARTMENTA | |
|---|---|---|--|
| Type of Document | Standard Code | Type of Document | Standard Code |
| Quality Manual Quality Procedure Work Instruction Form | TIEZA .QM.AA TIEZA.QP.CC TIEZA.WI.CC TIEZA.QF.CC | Quality Procedure Work Instruction Form | BBBB.QP.CC BBBB.WI.CC BBBB.QF.CC |

LEGEND:

- QM is Quality Manual
- QP is Quality Procedure
- WI is Work Instruction
- QF is Quality Form
- AA is the section number of the Quality Manual.
- BBBB is the designated code for each office (Sectoral/ Departmental).
- CC is the series number of the procedure, work instruction and form starting from 01.
- Includes Quality Workplace Procedures and Management Review.

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3.4.9 Office Number Code (Distribution List)/ Office Name and Office Code:

| Number Code OFFICE NAME | | OFFICE | |
|--|---|--------|--|
| 01 | Office of the Chief Operating Officer | 0000 | |
| 01A | Office of the Corporate Secretary | ocos | |
| 01B | Management Information Systems Department | MISD | |
| 01C | Internal Audit Department | IAUD | |
| 01D | Legal Department | LEGD | |
| 01E | Corporate Planning Department | COPD | |
| 01F | TIEZA Regulatory Office | TROF | |
| 02 | Office of the ACOO for Administration and Finance | AADF | |
| 02A | | | |
| 02B | 02B Financial Services Department | | |
| 02C | Travel Tax Department | FISD | |
| 03 | Office of the ACOO for Architectural & Engineering Services | AAES | |
| 03A | 3A Project Evaluation and Planning Department | | |
| 03B | Construction Management Department | COMD | |
| 04 | Office of the ACOO for Asset Management | AMGT | |
| 04A Operations Department | | OPED | |
| 04B Business Development Department | | BUDD | |
| 05 | Office of the ACOO for TEZ Management | ATEZ | |
| 05A | TEZ Regulation Department | TERD | |
| 05B TEZ Assistance and Monitoring Department | | TAMD | |

Example: OCOO. QP.01 (Quality Procedure number 01, originating from the Office of the Chief Operating Officer)

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3.5 Document Approval

Approval of Documents is in accordance to the following:

| Document | Poviewed L | | |
|---|-------------------------------|-------------------------|--|
| - countient | Reviewed by | Approved by | |
| Quality Manual | MANCOM | | |
| Quality Procedures | | Chief Operating Officer | |
| , | Quality Management | Chief Operating Office | |
| | Representative | | |
| Work Instruction | Sector Head | Quality Management | |
| | Offices under COO will be the | | |
| | Department Head | Representative | |

Approved by:

MARK T. LAPID

Chief Operating Officer

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