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09-01	Monitoring, Measurement, Analysis and Evaluation	3
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SUBJECT	FOREWORD		

Because of the highly competitive business environment, the adoption of Quality Management System is a strategic decision of any organization to promote the confidence of its business partners and the public. In view of this, the Department of Science and Technology (DOST) approved the implementation of DOST-GIA "Project 3: Establishment and Implementation of Quality Management System in Accordance with ISO 9001" under the DOST-GIA program "Establishment, Implementation and Maintenance of Management Systems in all DOST RDIs and Regional Offices."

The project is also in consonance with Executive Order No. 605, S. 2007 "Institutionalizing the Structure, Mechanisms and Standards to Implement the Government Quality Management Program, Amending for the Purpose Administrative Order No. 161, S. 2006", which was issued by then President Gloria Macapagal-Arroyo on 23 February 2007, directing all government agencies to adopt the ISO 9001:2000 Quality Management System as part of a government-wide quality management program.

Aligned with this initiative, the Industrial Technology and Development Institute (ITDI) has undertaken actions that help secure and strengthen its position as a service provider. The ITDI is *mandated* to conduct applied research and development on industrial manufacturing, minerals, chemicals, food processing and energy, packaging technology, biotechnology, transfer technologies and provide technical services and training.

Thus, to further ensure the continuous delivery of efficient and effective service, ITDI established, implemented and continuously maintain a quality management system in conformance with ISO 9001:2015 for the implementation of its technical operations.

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SUBJECT	OBJECTIVES OF THE QUALITY MANUAL		

The Quality Manual is documented with the following objectives:

- To provide consistent information, both internally and externally, about ITDI's Quality Management System;
- To demonstrate to the customers, suppliers, stakeholders and other interested parties, the ability of the ITDI to establish Quality Management System in meeting their requirements and expectations;
- To provide a documented system which serves as a basis in auditing the implementation and effectiveness of the QMS;
- To provide continuity of the company's QMS and its requirements during changing circumstances;
- To demonstrate compliance with the requirements of ISO 9001:2015 standard to which this QMS is based.

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	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 02-03
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SUBJECT	AUTHORIZATION FOR THE IMPLEMENTATION / UPDATING RESPONSIBILITY		

Implementation of the contents of the manual is authorized and approved by the Director effective on the date specified in the manual.

Updating of the manual is the responsibility of the Quality Management Representative following PM-QMS-07-01 Control of Maintain Documented Information.

Their specimen signatures appear below:

DR. ANNABELLÉ V. BRIONES, CESO IV

DR. ZORAYDA V. ANG
Quality Management Representative

ROCHEEL LEE C. DELUTA
Document Custodian

Prepared by:

Approved by:

DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 02-04
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SUBJECT	DISTRIBUTION OF THE (QUALITY MA	NUAL

The Quality Manual is distributed as follows:

COPY NO.	COPY HOLDER	DOCUMENT TYPE
Original	Document Custodian	PAPER
1	Director	-do-
2	Deputy Director for Research & Development	-do-
2A	Deputy Director for Administrative & Technical Services	-do-
3	Chief, ADM	-do-
4	Dropped (old copyholder Deputy QMR, R&D)	n/a
5	Chief, CED	-do-
6	Chief, EBD	-do-
7	Chief, FMD	-do-
8	Chief, FPD	-do-
9	Chief, MSD	-do-
10	Chief, PMISD	-do-
11	Chief, PTD	-do-
12	Chief, TSD	-do-
13	Head, Internal Audit Group	PAPER

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SUBJECT	CODING SYSTEM FOR	THE QUALIT	Y MANUAL

An alpha-numeric coding system is being followed in the Quality Manual as shown:

QM	ITDI	00 - 06	
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			Subject of the Manual
			Section of the Manual
	L		Name of Organization
			Type of Manual

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SUBJECT	DEFINITION OF TERMS		

The basis for the definition of terms is ISO 9001:2015 standard. The following abbreviations shall be used in this Manual.

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IDS -		Information and Documentation Section	
ICS -		Industrial Chemicals Section	
HRMS -		Human Resource Management Section	
FSS -		Food Safety Section	
FPD -		Food Processing Division	
FMD -		Finance and Management Division	
FES -		Food Engineering Section	
ESS -		Engineering Services Section	
ES -		Energy Section	
EBD -		Environment and Biotechnology Division	
DOST -		Department of Science and Technology	
DCI -		Document Custodian of the Institute	
DCD -		Document Custodian of the Division	
cs -		Cashier Section	
CPS -		Cleaner Production Section	
CED -		Chemicals and Energy Division	
BS -		Budget Section	
BDS -		Business Development Section	
BAC -		Bids and Awards Committee	
AS -		Accounting Section	
AMS -		Advanced Materials Section	
ADM -		Administrative Division	

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IFTS Industrial Fermentation Technology Section ITDI **Industrial Technology Development Institute** MIS **Management Information Systems** MSD **Materials Science Division** MTS **Microbial Treatment Section NCAR** Nonconformity and Corrective Action Report **PDLDS** Package Development & Label Design Section **PDS Product Development Section** PM **Procedures Manual PMISD** Planning and Management Information Systems Division **PMS** Planning and Monitoring Services **PPMS Property & Procurement Management Section** PS Pharmaceuticals Section PTD Packaging Technology Division **PTOSS** Package Testing and Other Services Section **QMR Quality Management Representative QMS Quality Management System RCTS Regional Cooperation and Training Section** RM**Risk Management RMS Records Management Section**

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/d/MR/	DIRECTOR

Surface Engineering Section

Special Materials Section

SES

SMS

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SUBJECT	DEFINITION OF TERMS		

TSD - Technological Services Division

WI - Work Instructions

WMS - Waste Management Section

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SUBJECT	COMPANY BACKGROUN	DINFORMA	TION

General Requirements

The ITDI through its *Director*, has established, documented, implemented and maintained a quality management system and continually improve its effectiveness in accordance with the requirements of ISO 9001:2015. The quality management system is designed following the concept of "Plan-Do-Check-Act" cycle.

The ITDI has identified the processes needed for its QMS and its application throughout the Institute and has determined the sequence and interaction of these processes. The criteria and methods needed to ensure the effectiveness of both the operation and control of these processes are established through the control parameters and performance measures integrated in the procedures of the QMS.

The Top Management has provided the necessary resources and information to support the control, operation and monitoring of these processes and related significant aspects. They ensure that processes within the scope of the QMS are monitored, measured and analyzed and actions necessary to achieve planned results and continual improvement of these processes are implemented.

Company Background Information

The Industrial Technology Development Institute (ITDI) is one of the seven research and development institutes (RDIs) under the Department of Science and Technology (DOST). The DOST-ITDI has been in existence since July 1, 1901 when it was then the Bureau of Government Laboratories (BGL) created through the Philippine Commission Act No. 156. It has undergone several reorganizations from 1905 until 1987, through a Republic Act (RA) and an Executive Order (EO). It is the only RDI among the other public sector research agencies in the country that is multidisciplinary, drawing on its expertise in the areas of chemicals, energy, biotechnology, environment, food processing, materials science, and packaging.

ITDI HISTORY

2009 – (Rationalization) August 26, 2009 – As a result of the Rationalization Program of the government under the Executive Order No. 366 dated 4 October 2004, the Industrial Technology Development Institute was rationalized with organizational and staffing modifications. Creation of the Packaging Technology Division and National Metrology Division was approved as well as consolidation of some divisions i.e. Fuels and Energy Division (FED) and Chemical and Mineral Division (CMD) to Chemicals and Energy Division (CED); Environmental Division (EnD) and Microbiology and Genetics Division (MGD) to Environment and Biotechnology Division (EBD); Economics and Planning Division (EPD) and Management Information Systems (MIS) to Planning and Management Information Systems Division (PMISD). RTID was likewise renamed to Technological Services Division.

1987 - The NSTA was reorganized into the Department of Science and Technology (DOST) by virtue of Executive Order Number 128 dated 30 January 1987.

Under this reorganization, NIST was renamed Industrial Technology Development Institute (ITDI) and remained one of the R&D institutes under the DOST. All centers were abolished and ITDI now has ten (10) technical divisions with (MSRI) now absorbed by ITDI. Seven (7) divisions to undertake

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R&D activities, three (3) to render technical services and two (2) support divisions were created.

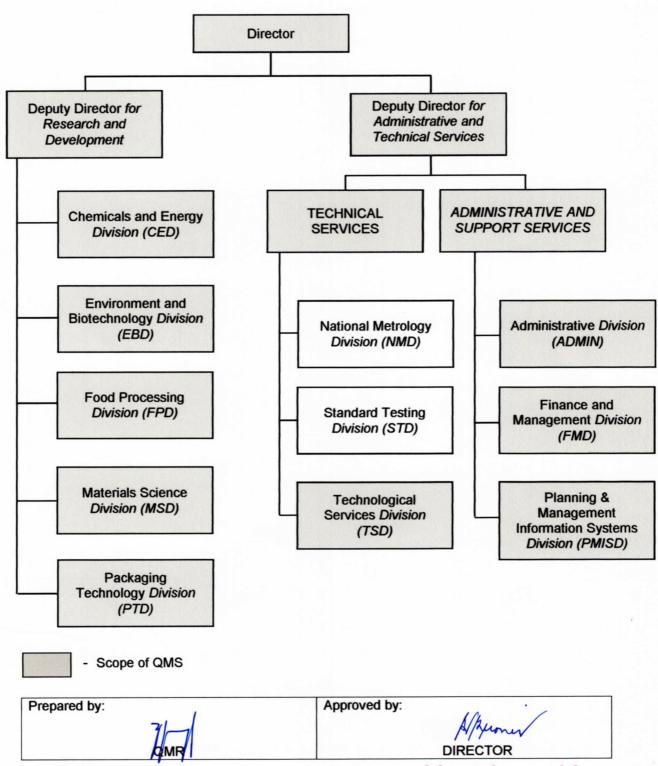
- 1982 By virtue of Executive Order Number 784 dated 17 March 1982, the NSDB was reorganized into the National Science and Technology Authority (NSTA). Under the reorganization NIST remained as one the R&D Institutes under the NSTA. As reorganized, research on agriculture was transferred to UPLB while the ceramics center emerged into an independent institute (MSRI). The Biological and Industrial research center were also abolished and in their place, three (3) centers were created namely: the National Research and Development (NRDC), Chemical Research and Development Center (CRDC), and the National Standards and Testing Center (NSTC). Under the NRDC and CRDC are seven(7) programs to undertake R&D activities, while the NSTC provides standardization and technical services.
- 1973 As part of the overall reorganization of the Executive branch of the government, the NIST was reorganized, but retained the same name. With the merger of the Agriculture Research Center, Biological Research Center and Medical Center, only two (2) technical R&D centers remained, namely Biological Research Center and Industrial Research Center. In addition, these were the Tests and Standards Laboratory and the Scientific Instrumentation Division to provide standardization and technical services.
- 1958 Under the so-called "Magna Carta of Philippine Science" RA 1067, NSB was reconstituted as the National Science Development Board (NSDB) which was designed to coordinate and supervise all scientific activities in the country. NSIRI became the National Institute of Science and Technology (NIST) under the supervision of NSDB.
- 1956 Congress approved RA Number 1606 authorizing the establishment of the National Science Board (NSB). IST was changed to the National Scientific and Industrial Research Institute (NSIRI), and was placed under supervision of NSB.
- 1951 The IS was renamed Institute of Science and Technology (IST) by virtue of Executive Order No. 392 and for the first time primarily concerned itself to industry-oriented research.
- 1947 The BS was transformed into the Institute of Science (IS) by virtue of Executive Order No. 94.
- 1934 The headship of the BS was passed on for the first time to a Filipino chemist. Dr. Angel S. Arguelles. The present-day Bureau of Soils, Bureau of Mines, Bureau of Fisheries and National Survey Division of Education Museum developed initially as part of the Bureau of Science during the pre-war years.
- 1905 By virtue of the Philippine Commission Act. No. 1407, the BGL was reorganized into the **Bureau** of **Science (BS)** and expanded its functions to include the Bureau of Mines and the Ethnological Survey Division of Education.
- 1901 1st of July The Bureau of Government Laboratories (BGL) came into existence through the Philippine Commission Act. No. 156. It was composed of the biological and chemical laboratories, a science library, and the Serum Laboratory of the Board of Health.

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Organizational Structure



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DOCUMENTED INFORMATION:

ISO 9001:2015 QMS Requirements

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The ITDI has identified the interested parties who are relevant to its quality management system and the requirements of the interested parties in order to prevent the potential effect on the organization's ability to consistently provide products and services, which meet the customer and applicable statutory and regulatory requirements.

Document Code	Process Title /		Description of Interested Parties		Their	Status of	Remarks
	Description		Internal	External	Requirements	Compliance	
PM-ADM- HRM-07-01	Human Resource Management	AOV, ADM- HRM	Employees Contract of Service Selection Board Staff Development Committee	CSC Ombudsman	Remuneration / Magna Carta Performance Evaluation Selection Criteria Contract Report	Complied	
PM-ADM- HRM-07-02	Staff Development	AOV, ADM- HRM	Employees	DOST	Training / Development	Complied	

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				CSC			
PM-ITDI-08- 01	Provision of Technical Services	Deputy Directors	Technical Staff	Academe Industry PNS Internal Standards ASTM BFAD DENR/DOE	Facilities Results of Test/Analysis Standardized Fees	Complied	
PM-ITDI-08- 02	Generation and Approval of Regular R&D Projects	Deputy Director, R&D	Researchers		Approved Project Proposal Budget Allocation	Complied	

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PM-ITDI-08-	Implementation	Deputy	Project Leader	DOST	Machine/Material	Complied	
03	of Regular Approved R&D	Director, R&D	PMISD	External Providers	Resources		
	Projects	NOLD	FIVII3D	External Providers	Report		
			R&D Chief	Fabricators			
	-		hard said to		Timely delivery of		
			BAC/PPMS	Academe	requisition items		
				Industry	Results of studies		
PM-CED-08-	Energy Audit /	Supv. SRS,	Energy Staff	DOE	Machine /	Complied	
01	Assessment	CED-ES	Division Chief	Industry	Material Resources		
				,	Mesour ces		
					Honorarium		
					Standardized Fees		
					EA Report		

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PM-EBD-	Environmental	Supv. SRS,	CPS Staff	DOST	Machine /	Complied	
CPS-08-01	Technology	EBD-CPS			Material		
	Verification		Section Head	DENR	Resources		
				Industry	Honorarium		
			-		Standardized Fees		
					ETV Report		
PM-PTD-08- 01	Label Design	Supv. SRS, PTD	Designer	ISTA	Packaging Technology	Complied	
01	Development	PID		IAPRI	Resources		
				IACM	Resources		
		_		FDA/USDA	Standardized Fees		
				Industry	Training &		
					Development		
					Design Folio		

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Approved Label Design Complied DTI PM-TSD-Supv. SRS, **BDS Staff** Technology TSD-BDS BDS-08-01 Transfer / Commercialization IPO of ITDI Matured Industry **Technologies** Machine / Supv., SRS, Writer's Pool Social Media PM-TSD-IDS-Creation and Material Committee TSD-IDS 08-01 Production of Academe Resources Social marketing TSD Chief Collaterals Training & Industry Development **Concerned Division** MOA/Contract Photographer Service **Layout Artist Published** Collaterals Complied **TSD Chief** Regional **Travel Order** Sr. SRS, TSD-PM-TSD-**Facilitation** Coordinator RCTS-08-01 **RCTS** Regional

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	Cooperation and Training		Training Coordinator Resource Person / Tech Staff	Customer /Participant	Schedule / Program of Activities CL/MOA Technology /	
PM-TSD- RCTS-08-02	Planning and Scheduling of Regular / Funded	Sr. SRS, TSD- RCTS	Section Head, TSD- RCTS	DOST Regional Offices	training Travel Order CL/MOA	
	Technology Transfer Program		Division Chief Technical Experts Director		Letter of Request	
PM-TSD- RCTS-08-03	Implementation of Technical and Other Regional Intervention	Sr. SRS, TSD- RCTS	Technology Expert RCTS Staff	Participant	Travel Order Technology	

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Honorarium CL/MOA Schedule / Program of Activities PM-TSD-Sr. SRS, TSD-Planning and **Division Chief** Academe Training Program/ RCTS-08-04 Scheduling of **RCTS** Module **Regular Training** Section Head, Industry **RCTS** Technology for program Commercialization Training Coordinator **Division Chief** PM-TSD-Training Sr. SRS, TSD-Training **Participant Training Program** Implementation RCTS-08-05 **RCTS** Coordinator / Manual **Training Facilitator** Honorarium

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3081501	PAR	TIES			

			Resource Person		Certificate of Completion Technology Innovation	
PM-TSD- RCTS-08-06	Design and Development of New Training Program	Sr. SRS, TSD- RCTS	Training Coordinator TSD Chief Design Team Resource Person		Course Design / Training Module Evaluation Criteria Certificate	
PM-PMISD- 08-01	Planning, Monitoring, and Evaluation of R&D Project	Supv. SRS, PMISD-PMS	Top Management PMISD R&D Division	DOST DBM COA	Approved Budget Accomplishment Report / Performance	

Prepared by:	Approved by:
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/∆ MR /	DIRECTOR

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CUDIFCT	UNDERSTANDING THE NEEDS AN	EXPECTATION	IS OF INTERESTED
SUBJECT	PAR	TIES	

	(Regular & Assisted)		TRC	Councils	R&D Programs and Projects Budget Utilization		
PM-FMD- AS-08-01	Billing	AO IV, FMD- BS	AO IV	DOST	Order of Payment	Complied	
			Chief, FMD	COA	Statement of Account		
			Accountant	DBM			
				Land Bank			
				GAM/NGAS			
				Bureau of Treasury			
				External Provider			
		J. January		Contractor			

Prepared by:	31-1	Approved by:	
	QMR/	DIRECTOR	

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				Fabricator		
ADM-PPMS- 08-01	Purchasing of Goods	BAC Secretariat	BAC / TWG	PhilGEPS	Specification requirement	Complied
			End-user	DBM-PS		
		AOV, ADM-			Manpower	
		PPMS	PPMS Staff	Eligible External Provider	complement	
	err binetie		Divisional		Budget allocation	
			Canvasser	Fabricator		
	1768 - 27 9 1				Plan / Design	
			Accounting /	Consultant		
	TOUR LINES	Albert 1	Budget Officer		Contract	Maria Maria W
	13 × 3 × 3 ×			Cooperative		
		3-22	Director / Deputy		Report	20 H 7 H L
			Director	COA		
				GPPB		
	for the second	. Patricker		NGAS / GAM		

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SUBJECT	UNDERSTANDING THE NEEDS AND EXPECTATIONS OF INTERESTED PARTIES				

PM-ADM- PPMS-08-02	AOV, ADM- PPMS	PPMS Staff	DBM-PS	On-time payment	Complied
		Inspection	COA	Facilities	
		Committee		(Laboratory /	
			Eligible External	Storage Room)	
		End-user	Provider		
				Report	

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SUBJECT	SCOPE OF THE QUALITY	MANAGEME	ENT SYSTEM

The Quality Management System of ITDI located at *the* DOST Compound, Gen. Santos Ave., Bicutan, Taguig City covers all processes under the Research and Development Divisions, Planning and Management Information Systems Division, Technological Services Division, and other support processes under the Finance and Management and Administrative Divisions.

The scope of certification is:

Generation and transfer of innovative technologies and provisions of scientific and technological services in the fields of biotechnology, chemicals and minerals, environment, foods processing, fuels and energy, materials science and packaging.

The processes involved are as follows:

RESEARCH AND DEVELOPMENT / TECHNICAL SERVICES PROCESSES

- Provision of Technical Services
- · Generation and Approval of Regular R&D Projects
- Implementation of Approved Regular R&D Projects

Chemicals and Energy Process

Energy Audit

Environment and Biotechnology Processes

- Cleaner Production Assessment
- Environmental Technology Verification
- Rapid Cleaner Production Assessment

Packaging Technology Process

Label Design Development

PLANNING AND MANAGEMENT INFORMATION PROCESSES

Planning, Monitoring and Evaluation of R&D Project

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QMR	DIRECTOR

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TECHNOLOGICAL SERVICES PROCESSES

- · Facilitating Regional Cooperation and Training
- Planning and Scheduling of Regular / Funded Technology Transfer Program
- Implementation of Technical and Other Regional Intervention
- Planning and Scheduling of Regular Training Program
- · Implementation of Training
- Design and Development of New Training Program
- Technology Transfer / Commercialization of ITDI Matured Technologies
- Creation and Production of Social Marketing Collaterals
- Preventive Maintenance
- Corrective Maintenance

ADMINISTRATIVE PROCESSES

- Human Resource Management
- Staff Development
- Purchasing of Goods
- Acceptance, Storage and Release of Purchased Items

FINANCE & MANAGEMENT PROCESSES

- Billing
- Budget Preparation

GENERAL QMS PROCESSES

- Management Review
- Handling Customer Complaints
- Customer Satisfaction Measurement
- Control and Monitoring of Customer Supplied Materials

PROCESSES REQUIRED BY ISO 9001:2015

- Control of Maintain Documented Information
- Control of Retain Documented Information
- Control of Maintain and Retain Documented Information in Electronic Media
- Internal Audit
- Control of Nonconforming Products
- Corrective Action
- Risk Management

Prepared by:	Approved by:
QMR/	DIRECTOR

August 2022

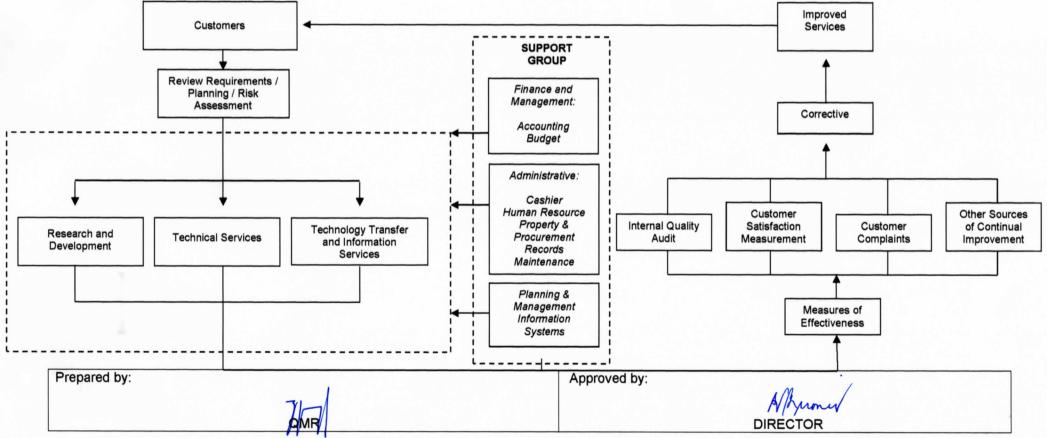
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Excluded from the Scope of Quality Management System are the 2 Technical Services Divisions: The Standard and Testing Division (STD, the National Metrology Laboratory (NML), and the Advance Device Materials Testing Laboratory. Most of the laboratories of the STD are already certified under ISO 17025, starting with the Microbiology Laboratory in 2002 and followed by the Organic & Inorganic Chemistry Laboratories, and the Physical and Performance Testing Laboratory. Likewise, the National Metrology Laboratory is also ISO 17025 certified by Deutsche Akkreditierungsstelle (DAkks) of Germany since 2010.

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SUBJECT	QUALITY MANAGEMENT SYSTEM AND ITS PROCESSES			

The QMS of ITDI covers processes that involve determining customer requirements, planning and managing the different activities involved to meet the requirements and expectations of its customers and conducting evaluation and improvement activities to enhance customer satisfaction. The diagram below illustrates the basic QMS processes.



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SUBJECT	LEADERSHIP AND COMMITMENT		

Top management has demonstrated leadership and commitment with respect to the quality management system by:

- Taking accountability for the effectiveness of the quality management system.
- Ensuring that the quality policy and quality objectives are established for the quality management system and are compatible with the context and strategic direction of the organization.
- Ensuring that integration of the quality management system requirements into the organization's business processes.
- Promoting the use of the process approach and risk-based thinking.
- Ensuring that the resources needed for the quality management system are available.
- Communicating the importance of effective quality management and of conforming to the quality management system requirements.
- Ensuring that the quality management system achieves its intended results.
- Engaging, directing and supporting persons to contribute to the effectiveness of the quality management system.
- Promoting improvement.
- Supporting other relevant management roles to demonstrate their leadership as it applies to their areas of responsibility.

Customer Focus

Top Management has demonstrated leadership and commitment with respect to customer focus by ensuring that:

- Customer and applicable statutory and regulatory requirements are determined, understood and consistently met.
- The risks and opportunities that can affect conformity or products and services and the ability to enhance customer satisfaction are determined and addressed.
- The focus on enhancing customer satisfaction is maintained.

Prepared by:	Approved by:
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DOCUMENTED INFORMATION:

Quality Policy
Quality Objectives
Functional Objectives
PM-QMS 09-03 Management Review
Minutes of Management Review
PM-QMS 09 -01 Customer Satisfaction Measurement

Prepared by:

Approved by:

DIRECTOR

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SUBJECT	QUALITY POLICY		

Establishing the Quality Policy

Top management has established, implemented and maintained a quality policy that:

- Is appropriate to the purpose and context of the organization and supports its strategic direction.
- Provides a framework for setting quality objectives.
- Includes a commitment to satisfy applicable requirements.
- Includes a commitment for continual improvements of the quality management system.

In pursuit of its commitment of providing customer satisfaction, Top Management has established a Quality Policy that reflects the values of employees of ITDI. The Quality Policy is stated as follows:

"We are committed to help local industries become globally competitive by providing appropriate technologies and services.

We shall continually improve our QMS to come up with conformity of products and services that would meet customer expectations within applicable regulatory and statutory requirements".

The quality objectives are:

- To complete and disseminate 90% of targeted R&D activities to meet the needs of local industries to become globally competitive.
- To deliver 90% of targeted technical services according to mutually agreed terms and conditions and existing applicable regulatory and statutory requirements.
- To attain at least a very satisfactory customer satisfaction rating for the year through the continual improvement of our QMS.

FUNCTIONAL OBJECTIVES:

Research & Development Group:

Chemicals Energy Division (CED)

- > To implement 90% of approved projects for implementation based on agreed duration.
- > To submit a terminal report two months after project completion.

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- > To implement 90% of the approved contract technical services / assistance based on agreed schedule with Very Satisfactory customer rating.
- > To undertake 90% of the approved technology transfer activities based on schedule with Very Satisfactory customer rating.

Environment & Technology Division (EBD)

- > To implement 90% of approved projects for implementation based on agreed duration.
- > To submit a terminal report two months after project completion.
- > To implement 90% of the approved contract and technical services based on agreed schedule with Very Satisfactory customer rating.
- > To undertake 90% of the approved technology transfer activities based on schedule with Very Satisfactory customer rating.

Foods Processing Division (FPD)

- > To implement 90% of approved projects for implementation based on agreed duration.
- > To submit a terminal report two months after project completion.
- > To implement 90% of the approved contract projects based on agreed schedule with Very Satisfactory customer rating.
- > To undertake 90% of the approved technology transfer activities based on schedule with Very Satisfactory customer rating.

Materials Science Division (MSD)

- > To implement 90% of approved projects for implementation based on agreed duration.
- > To submit a terminal report two months after project completion.
- > To implement 90% of the approved contract technical services / assistance on agreed schedule with Very Satisfactory customer rating.
- To undertake 90% of the approved technology transfer activities based on schedule with Very Satisfactory customer rating.

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Packaging Technology Division (PTD)

- > To implement on schedule 90% of the approved R&D projects and activities.
- > To submit a terminal report two months after the project completion.
- > To provide 90% of the request for packaging services with very satisfactory customer rating.

Support Group

Administrative Division - Cashier Section

 To ensure that 90% of approved disbursement vouchers are processed and transmitted to Agency's Government Servicing Bank within three (3) working days, subject to availability of funds.

Administrative Division - Human Resource Management Section

- To ensure that 90% of requisitioned employees are hired within the prescribed period in accordance to Omnibus Rules and Regulations.
- To ensure that at least one (1) Learning and Development is attended by every staff within a year based on available resources.
- To ensure that 100% of IPCR submissions are monitored and reviewed in accordance with SPMS calendar and analyzed to identify developmental interventions.

Administrative Division – Property & Procurement Management Section

- To ensure that 90% of approved Purchased / Work Orders are delivered within the agreed delivery terms based on the specifications of the end-user and in accordance with the provisions of RA 9184.
- To ensure that 90% of newly acquired, donation and transferred properties are properly accounted, labeled and recorded within fifteen (15) days upon inspection and acceptance of the end-user.
- To ensure that 100% of unserviceable properties are accounted and properly disposed within the scheduled plan in accordance with COA Rules and Regulations.

Prepared by:	Approved by:	* /
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SUBJECT	QUALITY POLICY		

Administrative Division - Records Management Section

- To ensure that 90% of records received are properly disseminated within the day and shall be maintained and readily available when needed.
- To ensure that 90% of records are controlled and disposed in accordance to provisions of RA 9470.

Finance & Management Division

- To ensure that all ITDI activities with budgetary requirements are acted upon within the prescribed period.
- To ensure that all financial transactions are processed within the prescribed period.
- To ensure that all budgetary request and financial transactions conform with accounting and auditing rules and regulations within the prescribed period.

Planning and Management Information Systems Division

- 1. To ensure the preparation of the Institute's overall plans and programs for implementation during the year.
 - > 90% of the Institute's programs and projects aligned *with* to the national and department priorities and needs.
 - > 90% of the Institute's plans and budget proposals prepared within the prescribed date.
- 2. To ensure Monitoring and Evaluation (M&E) of the Institute's programs, activities and projects (PAPs) during the year:
 - 2a. Monitoring
 - > 90% of projects monitored one month after the quarter.
 - 90% of projects monitored quarterly for risks identified.
 - > 90% of quarterly reports prepared.

2b. Evaluation

- 90% of project proposals evaluated with the ITDI-Technical Review Committee within the prescribed period.
- > 90% of ongoing projects evaluated with the ITDI-Technical Review Committee within two months after the quarter.
- 2c. Agency Performance Review
 - Preparation of the semestral Agency Performance Review Report two months after each semester with 100% accuracy.

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- 3. To ensure the review of existing and/or formulation of new organizational policies for improved operations and processes and effective governance with 100% acceptability.
- 4. To ensure the implementation of appropriate information systems for the projects, services and other activities of the Institute as identified in the approved Information Systems Strategic Plan (ISSP, 2020-2022) and in accordance with the approved budget allocation; and
- 5. To ensure action / response to requests for technical support on all ICT components:
 - 90% of requests for computer network technical support rendered within the agreed period.
 - 90% of requests for hardware technical support rendered with the agreed period.
 - 90% of requests for ICT System maintenance and support rendered within the agreed period.
 - 90% of requests for email technical support rendered within the agreed period.
 - 90% of requests for website maintenance and support rendered within the agreed period.

Technological Services Division

- To design and validate one new and acceptable training program within the year.
- To conduct 100% of the approved number of training programs for the year with a very satisfactory rating.
- To implement 100% of the approved requests for technical assistance / technology transfer in the Regions based on agreed schedules/duration within the year with a very satisfactory rating.
- To provide 100% relevant information to customers and stakeholders, both internally and externally in all forms (print and non-print, media / press conference, exhibits, techno fora) at the right time.
- To prepare techno-economic studies (market/industry profile, FS / BOP and techno-packages) of identified mature / commerciable technologies within the prescribed period.
- To develop business development strategies for marketing and promotion of identified technologies / new services within the prescribed period.
- To provide engineering related services in support of ITDI activities such as: R&D, technical services, maintenance and repair of physical facilities/buildings and technology transfer within the prescribed time with a very satisfactory rating.

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MR	DIRECTOR

August 2022

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SUBJECT	QUALITY POLICY		

Quality Management System:

Internal Quality Audit

- To ensure that internal audits shall be conducted once a year in accordance with ITDI's annual
 audit program. A detailed audit plan shall state the assignment of auditors in audit schedules as
 confirmed by auditees to determine whether the quality management system of ITDI:
 - Conforms to the planned arrangements of ISO 9001:2015; and
 - > Is implemented and maintained.
- To make sure that the audit shall be carried out in accordance with 9.2 of ISO 9001:2015, taking
 into consideration the status and importance of the processes in the areas to be audited, and the
 results of the previous audits with corrections and corrective actions.
- To render audit reports stating conformance or non-conformance to ISO 9001:2015, and recommendations to areas for improvement as inputs for management review. A follow-up of corrective and preventive actions for nonconformities, monitor and review of internal audit program shall be done in accordance with PM-QMS 09-02.

Document Custodian

- To ensure that 100% of ITDI QMS documents are updated, maintained and controlled in accordance with its effectivity date.
- To ensure that 100% of ITDI QMS documents are properly issued on its effectivity date and withdrawn from the copy holder at the right time.

DOCUMENTED INFORMATION:

PM-QMS 09-03 Management Review Minutes of Meeting

Prepared by:	Approved by:
James .	DIRECTOR

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SUBJECT	ORGANIZATIONAL ROLES, RESPONSIBILITIES AND AUTHORITIES		

Roles, Responsibilities and Authorities

The Top Management ensures that the responsibilities and authorities for relevant roles are assigned, communicated and understood within the organization through issuances of Office Orders, memoranda and during meetings in order to implement and maintain effective and efficient quality management system. The responsibilities and authorities of all employees throughout the organization are documented in the Manual of Actual Duties and Responsibilities. These are communicated to all employees during their orientation or their assignment to new positions and/or whenever there are changes to the position descriptions to enable them to contribute to the achievement of the quality objectives and to establish their involvement, motivation and commitment.

The Top Management assigns the responsibilities and authorities for:

- Ensuring that the quality management system conforms to the requirements of the International Standard.
- Ensuring that the processes are delivering their intended outputs.
- Reporting on the performance of the quality management system and addressing opportunities for improvement, in particular to top management.
- Ensuring the promotion of customer focus throughout the organization.
- Ensuring that the integrity of the quality management system is maintained when changes to the quality management system are planned and implemented.

The Director designated the Chief *Planning* Officer of ITDI to discharge the functions as the Quality Management Representative, on her behalf.

The responsibilities and authorities of the key personnel in the implementation and maintenance of the quality management system are described below:

<u>Director</u>

- · Formulates the organizational plans and methods for the internal operations of ITDI;
- Chairs the ISO management review meetings; and
- Provides the necessary resources for the effective implementation and maintenance of the quality management system.

ISO Technical Committee

- Acts as the policy-making body responsible for setting the direction and overseeing the implementation of QMS activities;
- Reviews the effectiveness and suitability of the Institute's QMS;
- Approves the QMS related final deliverables; and

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JOMP.	DIRECTOR

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CODOLOT	AUTHORITIES		

 Conducts ISO management reviews on such agenda as nonconformities, audit results, quality problems, performance indicators, corrective and preventive actions, and quality objectives.

Quality Management Representative (QMR):

Chief, Planning and Management Information Systems Division

ISO Risk Review Committee:

Chairperson – Quality Management Representative Co-Chairperson – Head, Internal Quality Audit Members – representative from ODD / CED / PMISD

ISO Risk Management Group:

Chairperson – Chief, Technological Services Division Co-Chairperson – Institute Document Custodian Members: Divisional Representatives

ISO Internal Auditors:

Chairperson – Supervising Science Research Specialist, CED Co-Chairperson – Science Research Specialist, PMISD Science Research Specialist, ODD Members – Divisional Representatives

ISO Document Custodians:

Chairperson – Administrative Officer V, ADM Members – Divisional Representatives

Quality Management Representative

- Ensures that the quality management system is established, implemented and maintained in accordance with ISO 9001:2015 standard;
- Responsible for reporting to and updating the top management on the performance of the quality management system for review, including recommendations for improvements;
- Ensures the promotion of awareness of customer requirements throughout the organization;
 and
- Acts as liaison with external parties on matters relating to quality management systems.

Risk Review Committee (RRC)

- Reviews and approves the existing divisional risk registers and the risk register crafted by the Risk Management Group
- Recommends changes, as needed, to the risk registers.
- Evaluates the effectiveness of the proposed changes to the risk registers.
- Monitors and evaluates the mitigating measures indicated by the Risk Management Group affecting the Institute's operational risks and strategies.
- Works with Risk Management Group in ensuring that the risks are properly considered and addressed.

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Risk Management Group

- Develops and oversees the institute risk management system;
- Ensures that the system remains effective; and
- Ensures that immediate corrective actions are taken to minimize the impact when risks occur.

Head Internal Audit Group

- Prepares annual audit program;
- · Ensures timely preparation of audit plan and audit checklist; and
- Implements the QMS internal audit based on the approved audit program.

Document Custodian

- Maintains and controls the QMS documents, including the revision, formal issuance, distribution and withdrawal;
- Prepares Master Lists of Controlled Documents, Externally-Sourced Documents and Quality Records; and
- Serves as secretariat during Management Review and QMS related meetings.

Division Chief

- · Takes charge of the administrative operations of the division;
- Assist the QMR in ensuring that processes within the scope of the QMS are established, implemented & maintained;
- · Report to the QMR on the performance of the QMS and any need for improvement;
- Assist the QMR in ensuring the promotion of awareness of customer requirements throughout the organization; and
- Ensure that all finished products conform to customer's specified requirements.

Section Chief

- Takes charge for the operations of the section;
- · Reports to the Division Chief the performance of the QMS and any need for improvement; and
- Assist the Division Chief in ensuring the promotion of awareness of customer requirements throughout the organization.

Human Resource Management Section

- · Takes charge for recruitment, selection and placement of qualified personnel;
- Ensures the continuous supply of competent people to man the processes within the scope of the QMS:
- Ensures the effectiveness of the human resource interventions conducted for all personnel within the scope of the QMS;
- Ensures that relevant documents and records of employees are updated and filed;
- Responsible for the promotion of the QMS in the Institute;
- Identifies training needs of the Institute's employees in relation to their roles in the effective implementation and maintenance of the QMS; and
- Ensures that critical personnel and members are appropriately trained and competent to perform assigned task.

Prepared by:	Approved by:
OMR	DIRECTOR

August 2022

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Property & Procurement Management Section

- Ensures that all purchases conform to the specified requirements and are acquired at the right time from the right source to the best advantage of the *Institute*; and
- Ensures that suppliers are competent to provide quality products & services to the Institutes.

Technological Services Division

- Ensures that scheduled preventive maintenance of all equipment is implemented accordingly;
 and
- Ensures the timely implementation of corrective maintenance.

Budget and Cashier Section

· Ensures that all finished job orders are properly billed and collected.

DOCUMETED INFORMATION:

Manual of Actual Duties and Responsibilities ITDI Special Order

Prepared by:	Approved by:
MR	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE		QM-ITDI 06-01
103	Ovelity Management Ovetem Henry	REVISION NUMBER	2
	Quality Management System Manual	PAGE NUMBER	1 of 2
OF OTION	DI ANNING	EFFECTIVITY DATE	01 June 2021
SECTION	PLANNING	DOCUMENT TYPE	PAPER / PDF FILE
CUBIECT	ACTIONS TO ADDRESS RISKS AND		
SUBJECT	OPPORTUNITIES		

The ITDI considered the issues referred to the organization and its context, the requirements referred to the needs and expectations of interested parties, and determined the risks and opportunities that need to be addressed to:

- Give assurance that the quality management system can achieve its intended results.
- Enhance desirable effects.
- Prevent, or reduce, undesirable effects.
- · Achieve improvement.

The ITDI has planned:

- a) Actions to address these risks and opportunities.
- b) How to:
 - Integrate and implement the actions into its quality management system processes.
 - · Evaluate the effectiveness of these actions.

Interested Parties	Requirements	Risks and Opportunities	Effectiveness of Actions
External Providers	~ Communication ~ Payment Terms ~ Timely delivery	~ Communication gap ~ Payment and delivery terms not specified ~ Delayed delivery	~ Communication like: email, phone, ~ Payment and Delivery terms mention in PO
Customer	~ Quality products and services ~ Timely delivery of product and services ~ Response to complaint. ~ Proper communication channel.	~ Loss of trust ~ Loss of customer ~ Improve process and enhance customer satisfaction	~ Defined in documented information and review in Management Review meetings.
Statutory and Regulatory Body	~ Complying with the statutory and regulatory requirements.	~ output is not acceptable ~ aligned to government regulations, legal requirements, etc	Defined in documented information and review in Management Review meetings.

Prepared by:	Approved by:
OMR	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 06-01
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SUBJECT	ACTIONS TO ADDRESS RISKS AND OPPORTUNITIES		AND
SUBJECT			

Employees	~ Remuneration	~ fair and just compensation and/or benefits.	~ compliance with the Magna Carta for S&T personnel.
	~ Training and development	~ Missed training opportunity due to limited budget	~ Defined in Staff development and review in Management Review meetings.
		~ technical expertise	

DOCUMENTED INFORMATION:

Risk Register
Quality Objectives
Functional Objectives
Minutes of Meeting
Minutes of Management Review

Prepared by:	Approved by:
QMR /	DIRECTOR

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10	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 06-02
(C)	Quality Management Cyctem Manual	REVISION NUMBER	2
	Quality Management System Manual	PAGE NUMBER	1 of 2
SECTION	N PLANNING	EFFECTIVITY DATE	01 June 2021
OLOTION	LAMMING	DOCUMENT TYPE	PAPER / PDF FILE
CLID IECT	QUALITY OBJECTIVES AND PLANNING TO		NING TO
SUBJECT	ACHIEVE THEM		

The ITDI has established quality objectives at relevant functions, levels and processes needed for the quality management system. The quality objective is stated as follows:

- 1. To complete and disseminate 90% of targeted R&D activities to meet the need of local industries to become globally competitive.
- 2. To deliver 90% of targeted technical services according to mutually agreed terms and conditions and existing applicable regulatory and statutory requirements.
- 3. To attain at least a very satisfactory customer satisfaction rating for the year through the continual improvement of our QMS.

The quality objective is:

- · Consistent with the quality policy.
- Measurable.
- Taking into account applicable requirements.
- · Relevant to conformity of products and services and to enhancement of customer satisfaction.
- Monitored.
- · Communicated.
- Updated as appropriate.

In order to achieve its quality objectives, ITDI determined:

- What will be done?
- What resources will be required?
- Who will be responsible?
- When it will be completed
- How the results will be evaluated?

Prepared by:	Approved by:
AMR.	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 06-02
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OLID IFOT	QUALITY OBJECTIVES AND PLANNING TO ACHIEVE THEM		NING TO
SUBJECT			

DOCUMENTED INFORMATION:

Quality Objectives Functional Objectives Monitoring of Objectives

Prepared by:	Approved by:	
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ØMR	DIRECTOR	

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 06-03
(C)	Overting Management Overten Manual	REVISION NUMBER	1
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CECTION	ECTION PLANNING	EFFECTIVITY DATE	03 January 2020
SECTION		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	PLANNING OF CHANGES		

The ITDI has determined the processes needed for the quality management system and their application throughout the Institute's Quality Management System and its Processes.

When there are changes needed in the quality management system, the ITDI ensures that such changes are carried out in a planned manner and considers the following:

- The purpose of the changes and their potential consequences.
- · The integrity of the quality management system.
- · The availability of resources.
- The allocation or reallocation of responsibilities and authorities.

DOCUMENTED INFORMATION:

QM-ITDI 04-04 Quality Management System and its Processes PM-ITDI 06-01 Change Management Process

Prepared by:	Reviewed / Approved by:
James 1	DIRECTOR

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	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-01
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SECTION	ECTION SUPPORT	EFFECTIVITY DATE	01 June 2021
SECTION	SUPPORT	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	RESOURCES		

General

The ITDI determines and provides the resources needed for the establishment, implementation, maintenance and continual improvement of the quality management system. Thus, the ITDI considers the capabilities of, and constraints on, existing internal resources and what needs to be obtained from external providers.

The Top Management ensures that the resources essential to the implementation of strategies and the achievement of the institute's objectives are identified and that the same are made available.

These decisions on the allocation of resources for operation and improvement of the QMS, enhancement of employees' competence through training, education and human resource intervention, the satisfaction of customers and other interested parties and planning for future resource needs are made during planning sessions. These resources may include people, infrastructure, work environment, information, suppliers and partners, and financial resources.

The Plans are reviewed regularly to ensure the efficiency and the effectiveness of resource allocation.

People

The ITDI ensures that personnel performing work affecting quality are competent on the basis of appropriate education, training and experience.

To assess, enhance and develop further their competence in providing quality products to the customers, the Administrative Division in coordination with the divisions concerned conducts annual training needs analysis based on competencies defined for each position. The *Human Resource Management Plan (HRDP)* is prepared based on the identified training needs.

In addition to the provision of training, other actions and interventions are undertaken to address identified *competency* gaps in competence. Training and/or other actions taken are designed to ensure that all employees are aware of the relevance and importance of their activities and their contribution to the attainment of the quality objectives.

After each training or completion of other actions taken, the Chief, Administrative Division evaluates their effectiveness in improving/enhancing competence of employees.

During the implementation of the QMS, all employees shall be appropriately provided with training and/or orientation to make them aware of the following:

- The importance of conformity with the quality policy and procedures and with the requirements of the quality management system;
- Their roles and responsibilities in achieving conformity with the requirements of the quality management system; and

The Records of education, training, skills and experience of employees are included in the 201 file.

Prepared by:	Approved by:
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QMR/	DIRECTOR

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SUBJECT	RESOURCES		

Infrastructure

The ITDI determines, provides, and maintains the environment necessary for the operations of its processes and to achieve conformity of products and services. The infrastructure includes building and associated utilities, equipment, including hardware and software, transportation resources and information and communication technology.

Environment for the Operation of Processes

To achieve conformity of products and services, *the* ITDI determines, provides and maintains a suitable environment which is combination of human and physical factors such as social, psychological, and physical environment.

The ITDI also implements program to ensure that work can be performed effectively, efficiently, and safely employees' opportunities for greater involvement thus, realizing the potential of people in the Institute.

Monitoring and Measuring Processes

General

ITDI determines and provides the resources needed to ensure valid and reliable results when monitoring or measuring is used to verify the conformity of products and services to requirements. It ensures that the resources provided are suitable for the specific type of monitoring and measurement activities being undertaken and are maintained to ensure their continuing fitness for their purpose.

The Institute retained documented information as evidence of fitness for purpose of the monitoring and measurement resources.

Measurement Traceability

To ensure valid results, the measuring instruments and equipment are calibrated or verified or both at specified intervals and adjusted or re-adjusted as necessary. The measuring equipment is identified to determine the calibration status, safeguarded from adjustments that would invalidate the measurement result and be protected from damage and deterioration during handling, maintenance and storage.

In order to provide confidence in data, *the* confirmation that the measuring devices are fit to use and are maintained to suitable accuracy and accepted standards, and identification of status are included in the measuring and monitoring processes.

When the equipment is found not to conform to requirements, the concerned division/section assesses and records the validity of the previous measuring results.

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August 2022
Approved by:
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N/Zmoner/
DIRECTOR

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SUBJECT	RESOURCES		

Organizational Knowledge

The Institute determines the knowledge necessary for the operation of its processes and to achieve conformity of products and services. This organizational knowledge based from internal and external sources are maintained and be made available to the extent necessary.

When addressing the needs and trends, the ITDI considers its current knowledge and determines how to acquire or access any necessary additional knowledge and required updates, to enhance the effectiveness of the Quality Management System and enhance customer satisfaction.

DOCUMENTED INFORMATION:

Minutes of Planning Sessions Minutes of Staff Meetings **Annual Training Plan** PM-ADM-HRM 07-01 Human Resource Management PM-ADM-HRM 07-02 Staff Development PM-ADM-PPMS-08-01 Purchasing of Goods Training Record 201 File **Preventive Maintenance Corrective Maintenance** 5S Program Calibration Plan of Inspection, Measuring and Test Equipment **Database of Intellectual Property Patents Certificates Terminal Reports Technical Papers Travel Reports** Proceedings / Minutes of Planning Workshops

Prepared by:	Approved by:
3 mal	DIRECTOR
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SUBJECT	COMPETENCE		

The ITDI has determined the necessary competence of person(s) doing work under its control that affects the performance and effectiveness of the quality management system. It ensured that these persons are competent on the basis of appropriate education, training or experience. Wherever applicable, the Institute takes action to acquire the necessary competence, and evaluate the effectiveness of the actions taken. These applicable actions can include the provision of training, mentoring or re-assignment of currently employed persons, or hiring or contracting of competent persons.

The Institute retained documented information as evidence of competence.

DOCUMENTED INFORMATION:

Individual Development Plan (IDP) Human Resource Management Plan

Prepared by:	Reviewed / Approved by:
TH	Afrana
/Q/MR/	DIRECTOR

DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE		DOCUMENT CODE	QM-ITDI 07-03
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SUBJECT	AWARENESS		

The ITDI ensures that persons doing work under the Institute control are aware of the quality policy, the relevant quality objectives, their contribution to the effectiveness of the quality management system requirements and implications of not conforming with the quality management system requirements.

DOCUMENTED INFORMATION:

Quality Policy Quality Objectives Functional Objectives.

Prepared by:	Approved by:
DIME	DIRECTOR

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	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-04
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	SUPPORT	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	COMMUNICATION		

The pertinent information related to the effective implementation of the QMS are relayed to all employees through Office Orders, memoranda, meetings and posting at strategic places.

The Internal communication to all levels and functions of the *Institute* shall include but not *be* limited to the following:

- · Quality Policy;
- · Quality objectives;
- · Procedures relevant to their duties and responsibilities; and
- Results of monitoring, measurement, and analysis needed to ensure effective implementation.

The Internal and external communications relevant to the quality management system are determined including: what it will communicate, when to communicate, with whom to communicate, how to communicate and who communicates.

INTERNAL COMMUNICATION

WHAT	WHEN	WITH WHOM	HOW	WHO
Quality Policy / Quality Objectives	Management Review	All Employees Interested Parties	Display / Paper / Electronic	Director / QMR / DC
Effectiveness of QMS (QM/PM/WI)	Training / Orientation	All Employees	Paper / Electronic	Director / QMR DC
Internal Audit Plan	Before the Audit (at least once a year)	Top Management Process Owners of all functional areas	Paper / Electronic	Internal Quality Auditors
Management Review	After the Internal Audit	ITDI EXECOM	Meeting (face-to- face/virtual)	Director
Strategic Plan	First Quarter Succeeding month after each quarter	Division Heads / Project Leaders / Project Staff	Letter / email / Virtual cascading / orientation	PMISD
Roles, Responsibilities and Authorities	During recruitment / Promotion / Department change	Employee	Procedure / Oral / Training	Director ADM-HRMS
ITDI Office Order	When necessary	All Employees	Letter / Meeting Email blast to all employees	Director

Prepared by:	Approved by:
OMR	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-04
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SUBJECT	COMMUNICATION		

Competency ~ Training needs ~ Training Plan	First quarter	EXECOM ITDI Staff	Paper / Electronic	Division Heads
Customer Complaint / Negative Feedback	When there are issues or rating is below VS.	Customer	Letter / Meeting	Process Owner / Division concerned / QMR / Director

EXTERNAL COMMUNICATION

WHAT	WHEN	WITH WHOM	HOW	WHO
Information to	During pre-	External	Purchase Order /	Property &
External	conference	Providers	Meeting /	Procurement
Providers	bidding /		Telephone /	Management
	Submission of		Website /	Section /
	Quotation		Email	Bids & Awards
				Committee /
				PMISD-
				Management
				Information
				System
Products and	During inquiry	Customer	Letter / email	Division Chief /
Services	thru email / social		Meeting	Project Leader /
Information	media platform		MOA /	Staff
Incoming and	When necessary	All Employees /	Letter / email	Process Owner
Outgoing		Interested parties		
Correspondences				
Customer	When there are	Customer	Letter / Meeting	Process Owner /
Feedback / Action	issues or rating is			Division
Taken for	below VS.			concerned /
Customer				QMR /
Complaints	2			Director

DOCUMENTED INFORMATION:

Quality Policy
Quality Objectives
Functional Objectives
Office Orders
Incoming/Outgoing Correspondences
Minutes of Meeting
Records of Orientation/Re-orientation

Prepared by:	Approved by:
7//	Approved
ø i nr/	DIRECTOR

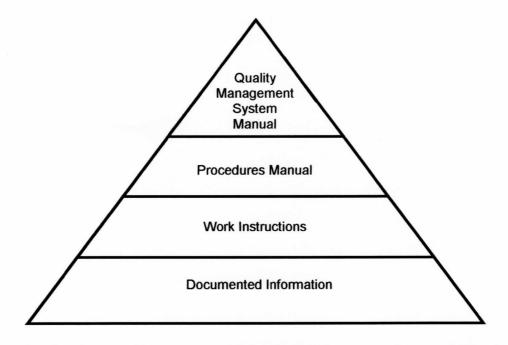
	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-05
SECTION	QUALITY MANUAL QUALITY MANAGEMENT SYSTEM REQUIREMENTS	REVISION NUMBER	3
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		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	DOCUMENTED INFORMATION		

The documentation of ITDI QMS is embodied in the documented statements of quality policy, objectives, quality manual, the mandatory *processes* required by ISO 9001:2015 and other documents needed by ITDI to ensure the effective planning, operation and control of the identified processes.

The Quality Management System Manual includes:

- The scope of the QMS as well as the details and justification for the exclusions in the QMS requirements;
- · The linkage of the procedures to the QMS; and
- The description of the interaction between the processes of the QMS.

The hierarchy of the QMS documentation is shown below:



The required "maintain and retain documented information" is complied as evidence of implementation.

The QMR is responsible for the implementation of this policy.

Prepared by:	Approved by:
QMR/	DIRECTOR

1 0	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-05
SECTION	QUALITY MANUAL QUALITY MANAGEMENT SYSTEM	REVISION NUMBER	3
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SECTION	REQUIREMENTS	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	DOCUMENTED INFORMATION		

Creating and Updating

The ITDI ensures that internally-generated and documents of external origin (in both soft and hard copies) needed for the effective implementations of its QMS are controlled. A documented procedure is maintained by the Document Custodian of the Institute to ensure appropriate:

- Identification and description (e.g.a title, date, author, or reference number).
- Format (e.g. language, software version, graphics) and media (e.g. paper, electronic).
- Documents are reviewed and approved for suitability and adequacy.
- A Master List of Controlled Maintain Documented Information is maintained by DCI and DCD to prevent the use of obsolete documents. The list includes documents needed for knowledgepreservation and legal purposes. Such documents are properly marked.

Control of Documented Information

The ITDI maintains and retains documented information to provide evidence of conformance to ISO 9001:2015 requirements and implementation of the QMS. All documented information required by the quality management system and by *the* International Standard are controlled to ensure:

- It is available and suitable for use where and when it is needed.
- It is adequately protected (e.g. from loss of confidentiality, improper use, or loss of integrity).

For the control of documented information, the ITDI addressed the following:

- Distribution, access, retrieval and use.
- · Storage and preservation, including preservation of legibility.
- Control of Changes (e.g. version control).
- Retention and disposition.

A Master List of *Retained* Documented Information is maintained to ensure control. Maintenance and storage of confidential records are the responsibility of the concerned heads.

The documented information of external origin determined by the organization to be necessary for the planning and operation of the quality management system is identified as appropriate, and to be controlled.

Prepared by:	Approved by:
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ØMR/	DIRECTOR

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SUBJECT	DOCUMENTED INFORMATION		

The Documented information retained as evidence of conformity is protected from unintended alterations.

DOCUMENTED INFORMATION:

PM-QMS 07-01 Control of *Maintain*ed Documented Information
PM-QMS 07-02 Control of *Retain*ed Documented Information
PM-QMS 07-03 Control of *Maintained* and Retain Documented Information in Electronic Media
Master List of Controlled *Maintain*ed Documented Information
Master List of Quality *Retained* Documented Information
Master List of Externally-Sourced Documented Information

Prepared by:	Approved by:
31-1	Alproner
ØMR/	DIRECTOR

SECTION	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-06
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		DOCUMENT TYPE	PDF / PAPER
SUBJECT CONTROL OF MONITORING AND MEASURING EQUIPMENT			

The ITDI has determined the monitoring and measuring equipment needed and measurements to be undertaken to provide evidence of conformity of products to specified requirements. The processes to ensure that monitoring and measurement can be carried out and are carried out in a manner that is consistent with the monitoring and measuring requirements are established and maintained.

To ensure valid results, the measuring instruments and equipment are calibrated or verified or both at specified intervals and adjusted or re-adjusted as necessary. The measuring equipment is identified to determine the calibration status, safeguarded from adjustments that would invalidate the measurement result and be protected from damage and deterioration during handling, maintenance and storage.

In order to provide confidence in data, *the* confirmation that the measuring devices are fit to use and are maintained to suitable accuracy and accepted standards, and identification of status are included in the measuring and monitoring processes.

When the equipment is found not to conform to requirements, the concerned division/section assesses and records the validity of the previous measuring results.

The records of the results of calibration and verification are maintained.

DOCUMENTED INFORMATION:

Calibration Plan of Inspection, Measuring and Test Equipment

Prepared by:	Approved by:
7/-/	NAmores
(QMR/)	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 08-01
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SECTION	OPERATION	EFFECTIVITY DATE	01 June 2021
		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	OPERATIONAL PLANNING AND CONTROL		

The ITDI has planned, implemented and controlled the processes needed to meet the requirements for the provision of products and services, and implemented the actions determined in planning requirements for the products and services; established criteria for the processes and acceptance of products and services; determined the resources needed to achieve conformity to the product and service requirement; implemented control of the processes in accordance with the criteria; and determined, maintained and retained documented information to the extent necessary to have confidence that the processes have been carried out as planned and demonstrated the conformity of products and services to the customer's requirements.

The output of this planning is suitable for the Institute's operations.

The ITDI has controlled changes of the plan, reviewed the consequences of unintended changes, took action to mitigate any adverse effects, as necessary. It ensured that outsourced processes are controlled.

DOCUMENTED INFORMATION:

PM-ITDI 08-01 Provision of Technical Services

PM-ITDI 08-02 Generation and Approval of Regular R&D Projects

PM-ITDI 08-03 Implementation of Approved Regular R&D Projects

PM-CED 08-01 Energy Audit

PM-EBD-CPS 08-01 Cleaner Production Assessment

PM-EBD-CPS 08-02 Environmental Technology Verification

PM-EBD-CPS 08-03 Rapid Cleaner Production Assessment

PM-PTD 08-01 Label Design Development

PM-PMISD 08-01 Planning, Monitoring and Evaluation of R&D Project

PM-TSD-BDS 08-01 Technology Transfer / Commercialization of ITDI Matured Technologies

PM-TSD-IDS 08-01 Creation and Production of Social Marketing Collaterals

PM-TSD-RCTS 08-01 Facilitating Regional Cooperation and Training

PM-TSD-RCTS 08-02 Planning and Scheduling of Regular / Funded Technology Transfer Program

PM-TSD-RCTS 08-03 Implementation of Technical and Other Regional Intervention

PM-TSD-RCTS 08-04 Planning and Scheduling of Regular Training Program

PM-TSD-RCTS 08-05 Implementation of Training

PM-TSD-RCTS 08-06 Design and Development of New Training Program

PM-ADM-PPMS 08-01 Purchasing of Goods

PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased Items

PM-FMD-AS 08-01 Billing

PM-FMD-BS 08-01 Budget Preparation

Prepared by:	Approved by:
James 1	DIRECTOR

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SUBJECT	REQUIREMENTS FOR PRO	DUCTS AND	SERVICES

Customer Communication

The communication with customers include:

- Providing information relating to products and services.
- · Handling enquiries, contracts or orders, including changes.
- Obtaining customer feedback relating to products and services, including customer complaints.
- · Handling or controlling customer property.
- Establishing specific requirements for contingency actions, when relevant.

Determining the Requirements for Products and Services

When determining the requirements for the products and services to be offered to customer, the Institute ensures that:

- The requirements for the products and services are defined including any applicable statutory and regulatory requirements and those considered necessary by the organization.
- The Institute can meet the claims for the products and services it offers.

Review of the Requirements for Products and Services

The ITDI ensures that it has the ability to meet the requirements for products and services to be offered to customers. It conducts a review before committing to supply products and services to a customer, to include:

- Requirements specified by the customer, including the requirements for delivery and postdelivery activities.
- Requirements not stated by the customer, but necessary for the specified or intended use, when known.
- Requirements specified by the Institute.
- Statutory and regulatory requirements applicable to the products and services.
- Contract or order requirements differing from those previously expresses.

Prepared by:	Approved by:
DMR	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 08-02
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		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	REQUIREMENTS FOR PRO	DUCTS AND	SERVICES

The ITDI ensures that contract or order requirements differing from those previously defined are resolved. The customers' requirements are confirmed by the Institute before acceptance, when the customers do not provide a documented statement of their requirements.

The Institute retains documented information, as applicable:

- On the results of the review
- On any new requirements for the products and services

Changes to Requirements for Products and Services

The ITDI ensures that relevant documented information is amended, and that relevant persons are made aware of the changed requirements, when the requirements for products and services are changed.

DOCUMENTED INFORMATION:

PM-ITDI 08-01 Provision of Technical Services PM-ITDI 08-02 Generation and Approval of Regular R&D Projects

PM-ITDI 08-03 Implementation of Approved Regular R&D Projects

PM-CED 08-01 Energy Audit

PM-EBD-CPS 08-01 Cleaner Production Assessment

PM-EBD-CPS 08-02 Environmental Technology Verification

PM-EBD-CPS 08-03 Rapid Cleaner Production Assessment

PM-PTD 08-01 Label Design Development

PM-PMISD 08-01 Planning, Monitoring and Evaluation of R&D Project

PM-TSD-BDS 08-01 Technology Transfer / Commercialization of ITDI Matured Technologies

PM-TSD-IDS 08-01 Creation and Production of Social Marketing Collaterals

PM-TSD-RCTS 08-01 Facilitating Regional Cooperation and Training

PM-TSD-RCTS 08-02 Planning and Scheduling of Regular / Funded Technology Transfer Program

PM-TSD-RCTS 08-03 Implementation of Technical and Other Regional Intervention

PM-TSD-RCTS 08-04 Planning and Scheduling of Regular Training Program

PM-TSD-RCTS 08-05 Implementation of Training

PM-TSD-RCTS 08-06 Design and Development of New Training Program

PM-ADM-PPMS 08-01 Purchasing of Goods

PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased Items

PM-FMD-AS 08-01 Billing

PM-FMD-BS 08-01 Budget Preparation

Prepared by:	Approved by:	• ,
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	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 08-03
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OFOTION	OPERATION	EFFECTIVITY DATE	01 June 2021
SECTION	OPERATION	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	DESIGN AND DEVELOPMENT OF PRODUCTS AND SERVICES		

General

The ITDI has established, implemented and maintained a design and development process that is appropriate to ensure the subsequent provision of products and services.

Design and Development Planning

The Top Management ensures that the Research and Development Divisions such as CED, EBD, FPD, MSD, PTD, and concerned section of TSD of ITDI have determined the stages and controls for design and development.

For every R&D project and packaging design, the Project Team considers the following:

- The nature, duration and complexity of the design and development activities.
- The required process stages, including applicable design and development reviews.
- The required design and development verification and validation activities.
- The responsibilities and authorities involved in the design and development process.
- The internal and external resources needs for the design and development of products and services.
- The need to control interfaces between persons involved in the design and development process.
- The need for involvement of customers and user in the design and development process.
- The requirement for subsequent provision of products and services.
- The level of control expected for the design and development process by customers and other relevant interested parties.
- The documented information needed to demonstrate that design and development requirements have been met.

Whenever a customer requires a new technical training program that is not included in the master list of training program, the *heads* of TSD and EBD/Design Team *prepare* a design and development plan. The plan describes the different activities and defines responsibility for their implementation. It also describes the training objectives and expected outputs, sequencing of the design and development process, design methodologies, review, verification and validation activities. The plan is updated whenever changes in the design occur.

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SUBJECT	DESIGN AND DEVELOPMENT OF PRODUCTS AND SERVICES		

Design and Development Inputs

The inputs to design and development include functional and performance requirements, information derived from previous similar design and development activities, statutory and regulatory requirements, standards or codes of practice that the Institute has committed to implement, potential consequences of failure due to the nature of the products and services.

The inputs are adequate for design and development purposes, complete and unambiguous. The conflicting design and development inputs resolved.

The Concerned sections of CED, EBD, FPD, MSD, PTD, and TSD ensure that all inputs are reviewed for adequacy and that requirements are clear, unambiguous and not in conflict with each other.

The Institute has retained documented information on design and development inputs.

Design and Development Controls

The Institute applies controls to the design and development process to ensure that:

- · The results to be achieved are defined.
- Reviews are conducted to evaluate the ability of the results of design and development to meet requirements.
- Verification activities are conducted to ensure that the design and development outputs meet the input requirements.
- Validation activities are conducted to ensure that the resulting products and services meet the requirement
- Any necessary actions are taken on problems determined during the review, or verification and validation activities.
- Documented information of these activities is retained.

Design and Development Outputs

The CED, EBD, FPD, MSD, PTD, and TSD Design Teams ensure that the outputs of design and development meet the input requirements; adequate for the subsequent processes for the provision of products and services; include or reference monitoring and measuring requirements, as appropriate, and acceptance criteria; and specify the characteristics of the products and services that are essential for their intended purpose and their safe and proper provision.

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The Institute has retained documented information on design and development outputs. *The design outputs include* data demonstrating the comparison of process inputs to process outputs, product specifications including acceptance criteria, process, material and testing specifications, training requirements, user information, purchase requirements, and qualification tests and reports.

The design outputs are reviewed against inputs to provide objective evidence that outputs have effectively and efficiently met requirements for the process and product.

Design and Development Changes

The design and development changes are identified, reviewed and control of changes made during, or subsequent to, the design and development of products and services, to the extent necessary to ensure that there is no adverse impact on conformity to requirements.

The documented information are retained on: design and development changes; results of reviews; authorization of the changes; and actions taken, to prevent adverse impacts.

DOCUMENTED INFORMATION:

Design and Development Plan
Design and Development Inputs
Design and Development Controls
Design and Development Changes
Results of Reviews
Authorization of Changes
Actions Taken to Prevent Adverse Impacts

Prepared by:	Approved by:
MR	DIRECTOR

August 2022

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 08-04
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		EFFECTIVITY DATE	01 June 2021
		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	CONTROL OF EXTERNALLY PROVIDED PROCESSES, PRODUCTS AND SERVICES		

General

To ensure that externally provided processes, products and services conform to requirement, the Property & Procurement Management Section has determined the controls to be applied to externally provided processes, products and services when:

- Products and services from external providers are intended for incorporation into the Institute's own products and services.
- Products and services are provided directly to the customer(s) by external providers on behalf
 of the organization.
- A process, or part of a process, is provided by an external provider as a result of a decision by the Institute.

The Institute determines and applies the criteria for the evaluation, selection, monitoring of performance, and re-evaluation of external providers, based on their ability to provide processes or products and services in accordance with requirements

The Master List of Eligible External Providers is retained documented information and their performance is evaluated annually through the use of rating sheet.

Whenever purchase requests are made, *the* complete specification and description of items are indicated to ensure that only the required items are purchased. No purchases are made without the approval of the Director or her Authorized Representative. *The personnel* implementing the purchase are competent based on their education, training, and experience.

In case of subcontracted services for some processes that the organization's capability may not be adequate, the said services should be in conformance to the requirements of the customer.

Type and Extent of Control

To ensure that externally provided processes, products and services do not adversely affect the Institute's ability to consistently deliver conforming products and services to its customers, the Institute:

- Ensures that externally provided processes remain within the control of its quality management system.
- Defines both the controls that it intends to apply to an external provider and those it intends to apply to the resulting output.
- Takes into consideration the potential impact of the externally provided processes, products and services on the Institute's ability to consistently meet customer and

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applicable statutory and regulatory requirements and the effectiveness of the controls applied by the external provider.

 Determines the verification, or other activities, necessary to ensure that the externally provided processes, products and services meet requirements.

Information of External Providers

To ensure the adequacy of requirement prior to the communication to the external providers, the Institute communicates its requirement for:

- The processes, products and services to be provided.
- The approval of products and services, methods, processes and equipment, and the release of products and services.
- · Competence, including any required qualification of persons.
- The external providers' interactions with the organization.
- Control and monitoring of the external providers' performance to be applied by the Institute.
- Verification or validation activities that the organization, or its customer, intends to perform at the external providers' premises.

DOCUMENTED INFORMATION:

PM-ADM-PPMS 08-01 Purchasing of Goods PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased Items WI-ADM-PPMS 08-03 Supplier Performance Rating Master List of Eligible External Providers

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SECTION	Quality Management System Manual PRODUCT REALIZATION	REVISION NUMBER	4
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		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	SUBJECT PRODUCTION AND SERVICE PROVISION		

Control of Production and Service Provision

The ITDI implements production and service provision under controlled conditions that include, as applicable:

- The availability of documented information that defines: (1) the characteristics of the products to be produced, the services to be provided, or the activities to be performed; and (2) the results to be achieved.
- The availability and use of suitable monitoring and measuring resources.
- The implementation of monitoring and measurements activities at appropriate stages to verify that criteria for control of processes or outputs, and acceptance criteria for products and services, have been met.
- The use of suitable infrastructure and environment for the operation of processes.
- The appointment of competent persons, including any required qualification.
- The validation and periodic revalidation, of the ability to achieve planned results of the processes for production and service provision, where the resulting output cannot be verified by subsequent monitoring or measurement.
- The implementation of actions to prevent human error.
- The implementation of release, delivery and post-delivery activities.

Identification and Traceability

The Institute uses suitable means to identify outputs when it is necessary to ensure the conformity of products and services.

It identifies the status of outputs with respect to monitoring and measurement requirements throughout production and service provision.

The Provision of services is uniquely identified using Reference Number. The assigned Reference No. can be traced from the time the *customer*'s inquiry is received up to the time the product is released. The Reference/Project Number No. is indicated in all records.

The ITDI has retained the documented information necessary to enable traceability.

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Property Belonging to Customers or External Providers

The ITDI exercises care with materials, components, tools and equipment, premises, intellectual property and personal data belonging to customer or external providers while it is under the Institute's control or being used by the Institute. The Institute shall identify, verify, protect and safeguard customers' or external providers' property provided for use or incorporation into the products and services. If any customer's property is lost, damaged or otherwise found to be unsuitable for use, the Institute shall report this to the customer or external provider and retain documented information on what has occurred.

Preservation

The conformity of product during production and service provision and release to the intended destination are preserved by the ITDI in order to maintain conformity to requirements. The appropriate labels, handling instructions, contamination control, packaging, storage, transmission or transportation, and protection are ensured and protected to prevent damage during transport.

Post - Delivery Activities

To meet requirements for post-delivery activities associated with the products and services, the ITDI determines the extent of post-delivery activities that are required and considers:

- Statutory and regulatory requirements.
- The potential undesired consequences associated with its products and services.
- The nature, use and intended lifetime of its products and services.
- Customer requirements.
- Customer feedback.

Control of Changes

To ensure continuing conformity with requirements, the Institute reviews and controls changes for production or service provision, to the extent necessary to ensure continuing conformity with requirements.

The documented information describing the results of the review changes, the person(s) authorizing the change, and any necessary actions from the review are retained.

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DOCUMENTED INFORMATION:

PM-ITDI 08-01 Provision of Technical Services

PM-ITDI 08-02 Generation and Approval of Regular R&D Projects

PM-ITDI 08-03 Implementation of Approved Regular R&D Projects

PM-CED 08-01 Energy Audit

PM-EBD-CPS 08-01 Cleaner Production Assessment

PM-EBD-CPS 08-02 Environmental Technology Verification

PM-EBD-CPS 08-03 Rapid Cleaner Production Assessment

PM-PTD 08-01 Label Design Development

PM-PMISD 08-01 Planning, Monitoring and Evaluation of R&D Project

PM-TSD-BDS 08-01 Technology Transfer / Commercialization of ITDI Matured Technologies

PM-TSD-IDS 08-01 Creation and Production of Social Marketing Collaterals

PM-TSD-RCTS 08-01 Facilitating Regional Cooperation and Training

PM-TSD-RCTS 08-02 Planning and Scheduling of Regular / Funded Technology Transfer Program

PM-TSD-RCTS 08-03 Implementation of Technical and Other Regional Intervention

PM-TSD-RCTS 08-04 Planning and Scheduling of Regular Training Program

PM-TSD-RCTS 08-05 Implementation of Training

PM-TSD-RCTS 08-06 Design and Development of New Training Program

PM-ADM-PPMS 08-01 Purchasing of Goods

PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased Items

PM-FMD-AS 08-01 Billing

PM-FMD-BS 08-01 Budget Preparation

PM-QMS 08-01 Handling Customer Complaint

PM-QMS 08-02 Control and Monitoring of Customer Supplied Materials

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SECTION	OPERATION	EFFECTIVITY DATE	03 January 2020
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SUBJECT	RELEASE OF PRODUCTS AND SERVICES		

The ITDI implements planned arrangements, at appropriate stages, to verify that the product and service requirements have been met.

The release of products and services to the customer shall not proceed until the planned arrangements have been satisfactorily completed, unless otherwise approved by a relevant authority and, as applicable, by the customer.

The Institute retained documented information on the release of products and services that include:

- Evidence of conformity with the acceptable criteria.
- Traceability to the person(s) authorizing the release.

DOCUMENTED INFORMATION:

Records of Traceability / Logbook

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SECTION	OPERATION	EFFECTIVITY DATE	01 June 2021
		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	CONTROL OF NONCONFORMING OUTPUTS		

The ITDI ensures that product which does not conform to product requirements is identified and controlled to prevent its unintended use or delivery. It takes appropriate action based on the nature of the nonconformity and its effect on the conformity of products and services. It also applies to nonconforming products and services detected after delivery of products, during or after the provision of services.

The Institute deals with nonconforming outputs in one or more of the following ways:

- Correction
- Segregation, containment, return or suspension of provision of products and services.
- Informing the customer.
- Obtaining authorization for acceptance under concession.

The Top Management empowers the employees with the authority and responsibility to report nonconformities at any stage of a process in order to ensure timely detection and disposition of nonconformities. The authority for response to nonconformities and actions to be taken are defined in the documented procedures to maintain achievement of process and product requirements.

The use of, release and acceptance of nonconforming products is done under concession by divisions/sections chief concerned and, where applicable, by the customer.

All chiefs of divisions and sections concerned ensure that appropriate action be taken to preclude the original intended use or application of any nonconforming product.

Whenever applicable, all division chiefs and sections concerned ensure that appropriate action be taken to the effects or potential effects of the non-conformity when non-conforming product is detected after delivery or use has started.

The conformity to the requirements is verified when nonconforming outputs are corrected.

The Institute retained documented information that: described the nonconformity; describes the actions taken; describes any concessions obtained; and identifies the authority deciding the action in respect of the nonconformity.

DOCUMENTED INFORMATION:

PM-QMS 09-03 Control of Nonconforming Outputs

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01101507	MONITORING, MEASUREMENT, ANALYSIS AND			
SUBJECT	EVALUATION			

General

The Top Management ensures the effective and efficient measurement, collection and validation of data, conformity to product requirements as well as the customers' level of satisfaction. The validity and purpose of measurements and the intended use of data are reviewed to ensure that these processes add value to the Institute.

The performance of the Institute's processes includes measurement and evaluation of its services, capability of its processes, achievement of objectives, targets and programs, compliance to applicable legal and other requirements and satisfaction of customer.

The data derived from the monitoring of performance provide information and serves as guide in its continual improvement activities. The results of the analysis of data from improvement activities is considered as one input to management review in order to provide information for improving the performance of the Institute.

Applicable methods including statistical techniques and the extent of their use are included.

The ITDI evaluates the performance and the effectiveness of the quality management system. The QMR in consultation with all the respective divisional chiefs, implements the monitoring, measurement, analysis and improvement processes needed, as to:

- What needs to be monitored and measured.
- The method for monitoring, measurement, analysis and evaluation needed to ensure valid results.
- When the monitoring and measuring shall be performed.
- When the results from monitoring and measurement are analyses and evaluated.

The institute is retaining appropriate documented information as evidence of the results.

Customer Satisfaction

In order to measure the performance of the QMS, information relating to customer perception of the degree to which their needs and expectations have been fulfilled is monitored through the conduct of Customer Satisfaction Survey. The Institute determines the methods for obtaining, monitoring and reviewing this information.

The QMR in coordination with the Division Chiefs are responsible for Customer Satisfaction as one of the measurements of the Quality Management System and as to whether the Institute has met customer requirements. Customer satisfaction shall include collection of authentic data, frequency and validity of analysis. The trend and major elements affecting customer satisfaction and key indicators of customer dissatisfaction will be monitored supported by objective evidence.

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SUBJECT	MONITORING, MEASUREMENT, ANALYSIS AND EVALUATION		

A customer satisfaction survey is done *quarterly* and its analysis and presentation is done annually and presented in the Management Review. Any negative feedbacks are addressed immediately.

Analysis and Evaluation

The ITDI analyzes and evaluates appropriate data and information arising from monitoring and measurement. The results of analysis are used to evaluate:

- Conformity of products and services
- The degree of customer satisfaction.
- The performance and effectiveness of the quality management system.
- · If planning has been implemented effectively.
- The effectiveness of action taken to address risks and opportunities
- The performance of external providers.
- The need for improvements to the quality management system.

Monitoring and Measurement of Processes

The suitable methods for monitoring and, where applicable, measurement of the quality management system processes are applied to demonstrate the ability of the processes to achieve planned results. When planned results are not achieved, appropriate correction and corrective action are taken to ensure conformity of the product.

Monitoring and Measurement of Product

To verify that product requirements have been met, the Institute ensures that monitoring and measurement are carried out at appropriate stages of the product realization process in accordance with the planned arrangements.

An evidence of conformity with the acceptance criteria is maintained. The records indicate the person(s) authorizing the release of product. Only products which have been satisfactorily completed through the planned arrangements are released to the customer.

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SUBJECT	EVALUA	TION	

DOCUMENTED INFORMATION:

Accomplished Customer Feedback Survey Form Analysis of Data File/Report Plan for Preventive Action Action Plan Monitoring of Quality Objectives

Prepared by:	Approved by:
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SUBJECT	INTERNAL AUDIT		

Internal Audit

The Internal Audit Committee is responsible in planning and implementing internal audits in order to verify whether the quality management system conforms to the planned arrangements, to the requirements of ISO 9001:2015 and to the quality management system requirements established by the Institute and whether the QMS is effectively implemented and maintained.

An audit program is planned and conducted at least once a year, taking into consideration the status and importance of the processes and areas to be audited, as well as the results of previous audits. The Audit Plan defines the audit criteria, scope, frequency and methods of audit. The auditors are selected based on competence, training, work experience and qualification and shall not audit their own work. The objectivity and impartiality of the audit process are ensured in the selection of auditors and conduct of audits.

The responsibilities and requirements for planning and conducting audits, and for reporting results and maintaining records are defined in PM-QMS 09-02, Internal Audit.

The Heads responsible for *the* area being audited ensure that any necessary corrections and corrective actions are taken without undue delay to eliminate detected non-conformities and their causes.

The follow-up activities include the verification of the actions taken and the reporting of the verification results. The Internal audit reports include evidence of excellent performance in order to provide opportunities for recognition by management and motivation of people.

The Institute has retained documented information as evidence of the implementation of the audit programme and the audit results.

DOCUMENTED INFORMATION:

PM-QMS 09-02 Internal Audit Audit Program Audit Results

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SUBJECT	SUBJECT MANAGEMENT REVIEW		

The Top Management, together with the Quality Management Representative, concerned divisions and section heads review the effectiveness of the QMS at least once a year to ensure its continuing suitability, adequacy, effectiveness and alignment with the strategic direction of the organization. The review involves identifying opportunities for improvement and possible need for changes, including the quality policy and objectives.

The Management Review is planned and carried out taking into consideration the following inputs:

- a) The status of actions from previous management review.
- b) Changes in external an internal issues that are relevant to the quality management system.
- c) Information on the performance and effectiveness of the quality management system, including trends in:
 - 1. Customer satisfaction and feedback from relevant interested parties.
 - 2. The extent to which quality objectives have been met.
 - 3. Process performance and conformity of products and services.
 - 4. Nonconformities and corrective actions.
 - 5. Monitoring and measurement results.
 - 6. Audit results.
 - 7. The performance of external providers
- d) The adequacy of resources.
- e) The effectiveness of actions taken to address risks and opportunities.
- f) Opportunities for improvement.

The outputs of the *Management Review* include opportunities for improvement, any need for changes to the quality management system and resource needs.

Retain documented information from *Management Reviews* are maintained to provide for traceability and to facilitate evaluation of the *Management Review* process itself, to ensure its continued effectiveness and added value to the organization.

DOCUMENTED INFORMATION:

PM-QMS 09-01 Management Review

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SUBJECT	IMPROVEMENT		

General

The ITDI determines and selects opportunities for improvement and implements any necessary actions to meet customer requirements and enhance customer satisfaction, which include:

- Improve products and services to meet requirements as well as to address future needs and expectations.
- · Correct, prevent or reduce undesired effects.
- Improve the performance and effectiveness of the quality management system.

Non Conformity and Corrective Action

When a nonconformity occurs, including any arising from complaints, the QMR ensures that all nonconformities relating to product, process and quality system are investigated and the results are recorded. While investigating causes of non-conformances and deciding corrective actions, the QMR identifies other products / situations where same or similar nonconformance can occur.

The corrective actions for nonconforming products and processes, detected in the organization are recorded, analyzed, reviewed, and necessary actions are taken. This will be the responsibility of process owner. The corrective action and their effectiveness are monitored.

The documented procedures are established to define requirements for corrective action and for handling customer complaints.

The corrective actions are initiated whenever there are complaints from customers; audit findings; performance standards, objectives and targets are not met within the defined time frame; and nonconforming products, both internal and external to the organization occur.

The causes of nonconformities are analyzed and suitable action taken to prevent their reoccurrence. The effectiveness of the identified corrective action is monitored through customer feedback and also the customers are kept aware of the corrective actions initiated, with each identified corrective action.

Continual Improvement

The data generated from the monitoring and measurements of processes are analyzed regularly to determine areas that need to be continually improved to ensure effective implementation of QMS. This information is taken from the regular reviews of the quality policy, objectives, audit results, corrective and preventive actions, results of management reviews and analysis of customer feedback. The concerned heads are responsible for ensuring that continual improvement activities are carried out.

The documented procedure has defined the requirements for the determination of potential nonconformities and their causes, preparation and implementation of preventive actions where

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SUBJECT	IMPROVEMENT		

needed, and review effectiveness of actions taken. The Division Chiefs and QMR maintain records of actions taken and *are* presented during management review meetings.

DOCUMENTED INFORMATION:

PM-QMS 08-01 Handling Customer Complaint PM-QMS 09-04 Corrective Action

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	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT GODE	QM-ITDI 02-01
1	THE DESCRIPTION OF THE PARTY.	REVISION NUMBER	1:
	Quality Management System Manual	PAGE NUMBER	1 of 1
SECTION	USER'S GUIDE	EFFECTIVITY DATE	03 January 2020
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SUBJECT	FOREW	ORD	

Because of the highly competitive business environment, the adoption of Quality Management System is a strategic decision of any organization to promote the confidence of its business partners and the public. In view of this, the Department of Science and Technology (DOST) approved the implementation of DOST-GIA 'Project 3: Establishment and Implementation of Quality Management System in Accordance with ISO 9001" under the DOST-GIA program "Establishment Implementation and Maintenance of Management Systems in all DOST RDIs and Regional Offices."

Said project is also in consonance with Executive Order No. 605, S. 2007 "Institutionalizing the Structure, Mechanisms and Standards to Implement the Government Quality Management Program, Amending for the Purpose Administrative Order No. 161, S. 2006" which was recently issued by President Gloria Macapagal-Arroyo on 23 February 2007, directing all government agencies to adopt the ISO 9001:2000 Quality Management System as part of a government-wide quality management program.

Aligned with this initiative, the Industrial Technology and Development Institute (ITDI) has undertaken actions that help secure and strengthen its position as a service provider. The ITDI is given the mandate to conduct applied research and development on industrial manufacturing, minerals, chemicals, food processing and energy, packaging technology, biotechnology, transfer technologies and provide lechnical services and training.

Thus, to further ensure the continuous delivery of efficient and effective service, ITDI opted to establish, implement and maintain a quality management system in conformance with ISO 9001:2015 for the implementation of its technical operations.



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40	Quality Management System Manual	PAGE NUMBER	1 of 1
SECTION	USER'S GUIDE	EFFECTIVITY DATE	01 September 2016
		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	OBJECTIVES OF THE QU	JALITY MAN	IUAL

The Quality Manual is documented with the following objectives:

- To provide consistent information, both internally and externally, about ITDI's Quality Management System,
- To demonstrate to the customers, suppliers, stakeholders and other interested parties, the ability of the ITDI to establish Quality Management System in meeting their requirements and expectations;
- To provide a documented system which serves as a basis in auditing the implementation and effectiveness of the QMS;
- To provide continuity of the company's QMS and its requirements during changing circumstances;
- To demonstrate compliance with the requirements of ISO 9001:2015 standard to which this QMS is based.



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/ QMR	DIRECTOR	

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	DOCUMENT TYPE	PAPER / PDF FILE	
SUBJECT	AUTHORIZATION FOR THE IMPLEMENTATION UPDATING RESPONSIBILITY		March 1 - Table 1

implementation of the contents of the manual is authorized and approved by the Director effective on the date specified in the manual.

Updating of the manual is the responsibility of the Quality Management Representative following PM-QMS-07-01 Control of Maintain Documented Information

Their specimen signatures appear below:

DR. ANNABELLE V. BRIONES, CESO IV Director

Dr. DIANA L. IGNACIO Quality Management Representative

> ROCHEEL LEE C. DELUTA Document Custodian

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/ QMR	DIRECTOR

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		REVISION NUMBER	2
T	Quality Management System Manual	PAGE NUMBER	1 of 1
SECTION	USER'S GUIDE	EFFECTIVITY DATE	03 January 2020
		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	DISTRIBUTION OF THE	QUALITY MA	ANUAL

The Quality Manual is distributed as follows:

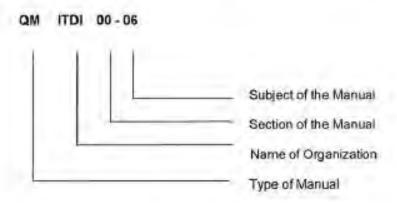
COPY NO.	COPY HOLDER	DOCUMENT TYPE
Original	Document Custodian	PAPER
1	Director	-do-
2	Deputy Director, R&D	-do-
3	Quality Management Representative/	-do-
4	Chief, CED	PDF File
5	Chief, EBD	-do-
6	Chief, FMD	-do-
7	Chief, FPD	-do-
8	Chief, MSD	-do-
9	Chief, PMISD	-do-
10	Chief, PTD	-da-
11	Chief, TSD	-do-
12	Head, Internal Audit Group	PAPER



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SECTION	USER'S GUIDE	EFFECTIVITY DATE	03 January 2020
		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	CODING SYSTEM FOR	THE QUALI	TY MANUAL

An alpha-numeric coding system is being followed in the Quality Manual as shown:





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	Quality Management System Manual	PAGE NUMBER	1 of 3
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		DOCUMENT TYPE	PAPER / PDF TYPE
SUBJECT	DEFINITION OF TERMS		

The basis for the definition of terms is ISO 9001:2015 standard. The following abbreviations shall be used in this Manual.

Prepared by:		Approved by:
IDS	-	
ics	-	Industrial Chemicals Section HNCONTDBLLED GOLY
HRMS	61	Human Resource Management Section
FSS	4	Food Safety Section
FPD	-	Food Processing Division
FMD	4	Finance and Management Division
FES	1	Food Engineering Section
ESS	-	Engineering Services Section
ES	1	Energy Section
EBD	2	Environment and Biotechnology Division
DOST	24	Department of Science and Technology
DCI		Document Custodian of the Institute
DCD	-	Document Custodian of the Division
CS	3	Cashier Section
CPS		Cleaner Production Section
CED	-	Chemicals and Energy Division
BS	4	Budget Section
BDS	-	Business Development Section
BAC		Bids and Awards Committee
AS	100	Accounting Section
AMS	-	Advanced Materials Section
ADM	1	Administrative Division

DIRECTOR

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SECTION	QUALITY MANAGEMENT SYSTEM	DOCUMENT TYPE	PAPER / PDF TYPE
SUBJECT	DEFINITION OF TERMS		

IFTS Industrial Fermentation Technology Section Industrial Technology Development Institute ITDI Management Information Systems MIS Materials Science Division MSD MTS Microbial Treatment Section Nonconformity and Corrective Action Report NCAR PDLDS Package Development & Label Design Section PDS Product Development Section PM. Procedures Manual Planning and Management Information Systems Division PMISD . PMS Planning and Monitoring Services PPMS Property & Procurement Management Section PS Pharmaceuticals Section Packaging Technology Division PTD PTOSS Package Testing and Other Services Section QMR Quality Management Representative QMS. Quality Management System RCTS Regional Cooperation and Training Section Risk Management RM Records Management Section RMS Surface Engineering Section SES

Prepared by:

SMS

duyw QMR Approved by:

Special Materials Section

DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 03-01
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SECTION	QUALITY MANAGEMENT SYSTEM	DOCUMENT TYPE	PAPER / PDF TYPE
SUBJECT	DEFINITION OF TERMS		

TSD - Technological Services Division

WI - Work Instructions

WMS - Waste Management Section



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SECTION	GOMBANN BROEKE	EFFECTIVITY DATE	03 January 2020	
	COMPANY PROFILE	DOCUMENT TYPE	PAPER / PDF FILE	
SUBJECT	COMPANY BACKGROUN	D INFORMA	TION	

General Requirements

ITDI through its Director has established, documented, implemented and maintained a quality management system and continually improve its effectiveness in accordance with the requirements of ISO 9001:2015. The quality management system is designed following the concept of "Plan-Do-Check-Act" cycle.

iTDI has identified the processes needed for their QMS and their application throughout the *Institute* and have determined the sequence and interaction of these processes. The criteria and methods needed to ensure the effectiveness of both the operation and control of these processes are established through the control parameters and performance measures integrated in the procedures of the QMS.

Top Management has provided resources and information necessary to support the control, operation and monitoring of these processes and related significant aspects

They ensure that processes within the scope of the QMS are monitored, measured and analyzed and actions necessary to achieved planned results and continual improvement of these processes are implemented.

Company Background Information

The Industrial Technology Development Institute or ITDI is one of the research and development institutes (RDIs) under the Department of Science and Technology. By virtue of Executive Order No. 128 dated January 30, 1987, ITDI is mandated to render a variety of services to local industries. It is the flagship agency of the Department, generating a large pool of technologies while providing technical services to industry.

ITDI HISTORY

2009 – (Rationalization) August 26, 2009 – As a result of the Rationalization Program of the government under the Executive Order No. 366 dated 4 October 2004, the Industrial Technology Development Institute was rationalized with organizational and staffing modification. Creation of the Packaging Technology Division and National Metrology Division was approved as well as consolidation of some divisions i.e. Fuels and Energy Division (FED) and Chemical and Mineral Division (CMD) to Chemicals and Energy Division (CED); Environmental Division (EnD) and Microbiology and Genetics Division (MGD) to Environment and Biotechnology Division (EBD); Economics and Planning Division (EPD) and Management Information Systems (MIS) to Planning and Management Information Systems Division (PMISD). RTID was likewise renamed to Technological Services Division.

1987 - The NSTA was reorganized into the Department of Science and Technology (DOST) by virtue of Executive Order Number 128 dated 30 January 1987.

Under this reorganization, NIST was renamed Industrial Technology Development Institute (ITDI) and remained one of the R&D institutes under the DOST. All centers were abolished and ITDI now has ten (10) technical divisions with (MSRI) now absorbed by ITDI. Seven (7) divisions to undertake R&D activities, three (3) to render technical services and two (2) support divisions were created.

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1982 - By virtue of Executive Order Number 784 dated 17 March 1982, the NSDB was reorganized into the National Science and Technology Authority (NSTA). Under the reorganization NIST remained as one the R&D Institutes under the NSTA. As reorganized, research on agriculture was transferred to UPLB while the ceramics center emerged into an independent institute (MSRI). The Biological and Industrial research center were also abolished and in their place, three (3) centers were created namely: the National Research and Development (NRDC), Chemical Research and Development Center (CRDC), and the National Standards and Testing Center (NSTC). Under the NRDC and CRDC are seven(7) programs to undertake R&D activities, while the NSTC provides standardization and technical services.

1973 - As part of the overall reorganization of the Executive branch of the government, the NIST was reorganized, but retained the same name. With the merger of the Agriculture Research Center, Biological Research Center and Medical Center, only two (2) technical R&D centers remained, namely Biological Research Center and Industrial Research Center. In addition, these were the Tests and Standards Laboratory and the Scientific Instrumentation Division to provide standardization and technical services.

1958 - Under the so-called "Magna Carta of Philippine Science" RA 1067, NSB was reconstituted as the National Science Development Board (NSDB) which was designed to coordinate and supervise all scientific activities in the country. NSIRI became the National Institute of Science and Technology (NIST) under the supervision of NSDB.

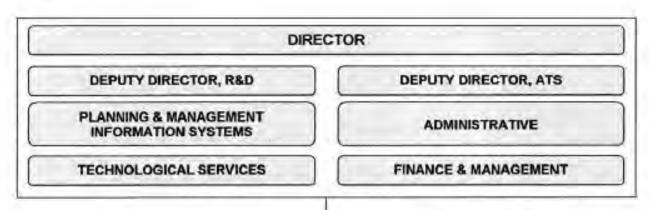
- 1956 Congress approved RA Number 1606 authorizing the establishment of the National Science Board (NSB), IST was changed to the National Scientific and Industrial Research Institute (NSIRI), and was placed under supervision of NSB.
- 1951 The IS was renamed Institute of Science and Technology (IST) by virtue of Executive Order No. 392 and for the first time primarily concerned itself to industry-oriented research.
- 1947 The BS was transformed into the Institute of Science (IS) by virtue of Executive Order No. 94.
- 1934 The headship of the BS was passed on for the first time to a Filipino chemist. Dr. Angel S. Arguelles. The present-day Bureau of Soils, Bureau of Mines, Bureau of Fisheries and National Survey Division of Education Museum developed initially as part of the Bureau of Science during the pre-war years.
- 1905 By virtue of the Philippine Commission Act. No. 1407, the BGL was reorganized into the Bureau of Science (BS) and expanded its functions to include the Bureau of Mines and the Ethnological Survey Division of Education.

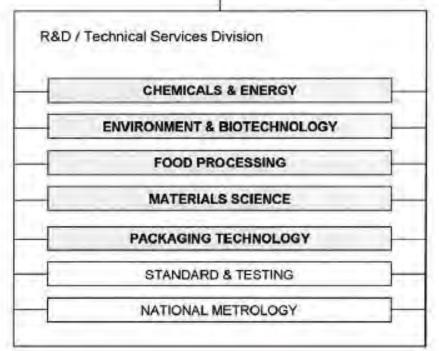
1901 - 1st of July - The Bureau of Government Laboratories (BGL) came into existence through the Philippine Commission Act. No. 156. It was composed of the biological and chemical laboratories, a science library, and the Serum Laboratory of the Board of Health.

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Organizational Structure





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COMPANY BACKGROUN	D INFORMA	TION
	Quality Management System Manual COMPANY PROFILE	Quality Management System Manual COMPANY PROFILE DOCUMENT CODE REVISION NUMBER PAGE NUMBER EFFECTIVITY DATE

DOCUMENTED INFORMATION:

ISO 9001:2015 QMS Requirements



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SUBJECT	UNDERSTANDING TH		ATION

ITDI determine external and internal issues that are relevant to its purpose and its strategic direction and that affect its ability to achieve the intended results of its quality management system through PEST (Political, Economic, Social and Technology factors) and SWOT (Strength, Weaknesses, Opportunities, and Threats) analysis. The external and internal issues identified through PEST and SWOT are continuously being monitored.

A. PEST

-	-	HARMAN	
POI	THE AL	VARIAR	1

- Changes in government leadership (local, national and international)
- Government policies to include trade regulation and requirements.
- International political situation.
- National and international policies and relationships.

ECONOMIC VARIABLES

- Membership to economic communities or organizations.
- 2. Government budget for STI.
- 3. Economic status / stability.
- 4. Changing industries trends.
- National and international economic agenda of the country.
- Globalization.
- Competitiveness amidst technological innovations
- Public good, participation in economic groups,
- Road map of other government agencies.

SOCIAL VARIABLES

- Philippine customer demographics (mostly female and young)
- Social media as a mode of communication.
- Cultural lifestyle
- Addressing the needs of vulnerable groups (victims of disaster and natural calamities)
- Foresight of other stakeholders and how to align with them.
- Sustainability of the current standing of the Institute and relationship with stakeholders.
- Crate avenues for potential scientists

TECHNOLOGICAL VARIABLES

- Internet of Things (IoT) connectivity.
- 2. Intellectual property protection.
- Technology gaps (local vis-à-vis international).
- Lifecycle of technologies.
- Timeliness of introduction of technology and innovation.
- Focus on innovation, including green innovation technologies, artificial intelligence, and industry 4.0.
- Setting up of new laboratories as indicated in the roadmap.

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Approved by

DIRECTOR

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B. SWOT

STRENGTHS

To meet the needs of local industries for global competitiveness:

- Multi-awarded experts and competent staff in the field of STI (local and international recognition).
- 2. High integrity personnel.
- ISO and internationally-recognized laboratories meeting regulatory and statutory requirements of customers.
- Multidisciplinary institute working as a cohesive team.
- Investments in modern R&D and technical services facilities.
- Active in dissemination of scientific and technical information.
- Strong resource generation capability.
- Institute with established national and international linkages.
- Competitive salary and other remunerations
- High customer satisfaction.

WEAKNESSES

- Need more industry collaboration to identify R&D needs
- Need to develop institutionalized system for impact assessment of technology and technical services disseminated to customers.
- 3. Low utilization of funds.
- Low level of customer awareness about ITDI and its capabilities.
- Low level of industry immersion of staff (researchers mobility)
- Need to intensify knowledge management (IP, success plan, data mining).

OPPORTUNITIES

- Change in political leadership allows refocusing Institute direction.
- New government policies and would help guide direction for R&D and technical services.
- Partnerships and collaborative R&D with international linkages to lessen technology gaps.
- R&D collaboration with local industries to address social and cultural issues.
- Additional avenue, e.g. Social media, internal connectivity, to promote ITDI and its services.
- Energy & Environment integration
- 7. Industry 4.0

THREATS

- Some facilities located on top of fault that compromising safety of staff and integrity of buildings and equipment.
- Government directive to streamline operations.
- Change in political leadership could disrupt ongoing activities.
- Other institutions compete with available limited funding.

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Approved by:

Approved by:

DIRECTOR

SUBJECT	UNDERSTANDING TH		ATION
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1000	Quality Management Protom Manual	REVISION NUMBER	3
	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 04-01

Documented Information:

Road Map ITDI Portfolio Strategic Plan



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	QMR	DIRECTOR

SUBJECT	UNDERSTANDING THE NEEDS AN	D EXPECTATION TIES	S OF INTERESTED
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ITDI has identified the interested parties who are relevant to its quality management system and the requirements of the interested parties in order to prevent the potential effect on the organization's ability to consistently provide products and services which meet the customer and applicable statutory and regulatory requirements.

Document	Process Title /	Process	Descripti	on of Intereste	d Parties	Their	Status of	Remarks
Code	Code Description Owner Client / External Stakeholders Providers	Regulatory Bodies	Requirements	Compliance				
PM-ADM- HRM-07-01	Human Resource Management	AOV, ADM- HRM	Employees Contract of Service Selection Board Staff Development Committee		CSC Ombudsman	Remuneration / Magna Carta Performance Evaluation Selection Criteria Contract Report	Complied	
PM-ADM- HRM-07-02	Staff Development	AOV, ADM- HRM	Employees	DOST	CSC	Training / Dev't	Complied	

SUBJECT	UNDERSTANDING THE NEEDS AN	D EXPECTATION	S OF INTERESTED
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Document Code	Process Title / Description	Process Owner	Description of Interested Parties			Their	Status of	Remarks
			Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
PM-ADM- PPMS-08- 02	Acceptance, Storage, and Release of Purchased Items	AOV, ADM- PPMS	PPMS Staff Inspection Committee End-user	DBM-PS Eligible External Provider	COA	On-time payment Facilities (Laboratory / Storage Room) Report		

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Approved by:

Approved by:

DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 04-02
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Document	Process Title /	Process	Description	on of Intereste	d Parties	Their	Status of	Remarks
Code	Description	Owner	Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
PM-ITDI- 08-01	Provision of Technical Services	Deputy Director, R&D	Technical Staff Academe Industry		PNS Internal Standards ASTM BFAD DENR/DOE	Facilities Result of Test/Analysis Standardize Fees	Complied	
PM-ITDI- 08-02	Generation and Approval of Regular R&D Projects	Deputy Director, R&D		9		Approved Project Proposal Budget Allocation	Complied	
PM-ITDI- 08-03	Implementation of Regular Approved R&D Projects	Deputy Director, R&D	Project Leader PMISD R&D Chief	DOST External Providers Fabricators		Machine/Material Resources Honorarium Report	Complied	rui

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SUBJECT	UNDERSTANDING THE NEEDS AN	D EXPECTATION	S OF INTERESTED	

PARTIES

Document	Process Title /	Process	Description	on of Intereste	d Parties	Their Requirements	Status of	Remarks
Code	Description	Owner	Client / Stakeholders	External Providers	Regulatory Bodies		Compliance	
			BAC/PPMS Academe Industry			Timely delivery of requisition items		
PM-CED- 08-01	Energy Audit / Assessment	Supv. SRS, CED-ES	Energy Staff Division Chief Industry		DOE	Machine / Material Resources Honorarium Standardized Fees EA Report	Complied	
PM-EBD- CPS-08-01	Environmental Technology Verification	Supv. SRS, EBD-CPS	CPS Staff Section Head Industry	DOST	DENR	Machine / Material Resources Honorarium Standardized Fees	Complied	TD

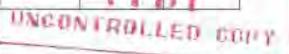
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	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 04-02

Document	Process Title /	Process	Description of Interested Parties			Their	Status of	Remarks
Code	Description	Owner	Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
						ETV Report		
PM-PTD- 08-01	Label Design Development	Supy. SRS, PTD	Designer	ISTA	FDA/USDA	Packaging Technology Resources Standardized Fees Training & Development Approved Label Design	Complied	
PM-TSD- BDS-08-01	Technology Transfer / Commercialization of ITDI Matured Technologies	Supv. SRS, TSD-BDS	BDS Staff Industry		DTI IPO		Complied	
PM-TSD- IDS-08-01	Creation and Production of Social marketing Collaterals	Supv., SRS, TSD-IDS	Writer's Paol Committee TSD Chief	Social Media		Machine / Material Resources Training & Development	CONTR	TU

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SUBJECT	UNDERSTANDING THE NEEDS AN	D EXPECTATION	S OF INTERESTED

UNDERSTANDING THE	NEEDS AND	EXPECTATIONS	OF INTERESTED
	PARTI	IES	

Document	Process Title /	Process	Description	on of Intereste	d Parties	Their	Status of	Remarks
Code	Description	Owner	Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
			Concerned Division Photographer Layout Artist Academe			MOA/Contract Service Published Collaterals		
PM TSD RCTS-08-01	Facilitation Regional Cooperation and Training	Sr. SRS, TSD-RCTS	TSD Chief Training Coordinator Resource Person / Tech Staff Regional Coordinator			Travel Order Schedule / Program of Activities CL/MOA	Complied	
			Customer /Participant				TTI	11

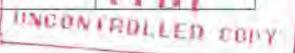


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SUBJECT

UNDERSTANDING THE NEEDS AND EXPECTATIONS OF INTERESTED PARTIES

Document	Process Title /	Process	Descripti	on of Intereste	d Parties	Their	Status of	Remarks
Code	Description	Owner	Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
PM-TSD- RCTS-08-02	Planning and Scheduling of Regular / Funded Technology Transfer Program	Sr. SRS, TSD-RCTS	Section Head, TSD-RCTS Division Chief Technical Experts Director	DOST Regional Offices		Travel Order CL/MOA Letter of Request TNA		
PM-TSD- RCTS-08-03	Implementation of Technical and Other Regional Intervention	Sr. SRS, TSD-RCTS	Technology Expert RCTS Staff Participant			Travel Order Honorarium CL/MOA Schedule / Program of Activities		
PM-TSD- RCTS-08-04	Planning and Scheduling of Regular Training program	Sr. SRS, TSD-RCTS	Division Chief Section Head, RCTS			Training Program/ Module Technology for Commercialization	TI	1



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Document	Process Title /	Process	Description	on of Intereste	d Parties	Their	Status of	Remarks
Code	Description	Description Owner	Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
			Training Coordinator Division Chief Academe Industry					
PM-TSD- RCTS-08-05	Training Implementation	Sr. SRS, TSD-RCTS	Training Coordinator Training Facilitator Resource Person Participant			Training Program / Manual Honorarium Certificate of Completion		
PM-TSD- RCTS-08-06	Design and Development of New Training Program	Sr. SRS, TSD-RCTS	Training Coordinator TSD Chief			Course Design / Training Module Evaluation Criteria	IT	li I

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Document	Process Title /	Process	Description of Interested Parties			Their	Status of	Remarks
Code	Description	Owner	Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
			Design Team Resource Person			Certificate		
PM-PMISD- 08-01	Planning, Monitoring, and Evaluation of R&D Project	Supv. SRS, PMISD-PMS	Top Management PMISD R&D Division TRC	DOST President's Agenda DBM	COA	Approved Budget Accomplishment Report R&D Programs and Projects		
PM-FMD- AS-08-01	Billing	AO IV, FMD-BS	AO IV Chief, FMD Accountant External Provider	DOST	COA DBM Land Bank GAM/NGAS	Order of Payment Statement of Account Magna Carta	Complied	

SUBJECT	UNDERSTANDING THE NEEDS AND EXPECTATIONS OF INTERESTED PARTIES		
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			Client / Stakeholders	External Providers	Regulatory Bodies	Requirements	Compliance	
			Contractor Fabricator		Bureau of Treasury			
PM-ADM- PPMS-08- 01	Purchasing of Goods	BAC Secretariat AOV, ADM- PPMS	BAC / TWG End-user PPMS Staff Divisional Canvasser Accounting / Budget Officer Director / Deputy Director	PhilGEPS DBM-PS Eligible External Provider Fabricator Consultant Cooperative	COA GPPB NGAS / GAM	Specification requirement Manpower complement Budget allocation Plan / Design Contract Report	Complied	



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SECTION	CONTEXT OF THE ORGANIZATION	DOCUMENT TYPE	PAPER / PDF FILE	
SUBJECT	SCOPE OF THE QUALITY I	MANAGEME	ENT SYSTEM	

The Quality Management System of ITDI located at DOST Compound, Gen. Santos Ave., Bicutan, Taguig City covers all processes under the Research and Development Divisions, Planning and Management Information Systems Division, Technological Services Division, and other support processes under the Finance and Management and Administrative Divisions.

The scope of certification is:

Generation and transfer of innovative technologies and provisions of scientific and technological services in the fields of biotechnology, chemicals and minerals, environment, foods processing, fuels and energy, materials science and packaging.

The processes involved are as follows:

RESEARCH AND DEVELOPMENT / TECHNICAL SERVICES PROCESSES

- Provision of Technical Services
- Generation and Approval of Regular R&D Projects
- Implementation of Approved Regular R&D Projects

Chemicals and Energy Process

Energy Audit

Environment and Biotechnology Processes

- Cleaner Production Assessment
- Environmental Technology Verification
- Rapid Cleaner Production Assessment

Packaging Technology Process

Label Design Development

PLANNING AND MANAGEMENT INFORMATION PROCESSES

Prepared by:

Approved by:

Approved by:

DIRECTOR

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TECHNOLOGICAL SERVICES PROCESSES

- Facilitating Regional Cooperation and Training
- Planning and Scheduling of Regular / Funded Technology Transfer Program
- Implementation of Technical and Other Regional Intervention
- Planning and Scheduling of Regular Training Program
- Implementation of Training
- Design and Development of New Training Program
- Technology Transfer / Commercialization of ITDI Matured Technologies
- Creation and Production of Social Marketing Collaterals
- Preventive Maintenance
- Corrective Maintenance

ADMINISTRATIVE PROCESSES

- Human Resource Management
- Staff Development
- Purchasing of Goods
- Acceptance, Storage and Release of Purchased Items

FINANCE & MANAGEMENT PROCESSES

- Billing
- Budget Preparation

GENERAL QMS PROCESSES

- Management Review
- Handling Customer Complaints
- Customer Satisfaction Measurement
- Control and Monitoring of Customer Supplied Materials

PROCESSES REQUIRED BY ISO 9001:2015

- Control of Maintain Documented Information
- Control of Retain Documented Information
- Control of Maintain and Retain Documented Information in Electronic Media
- Internal Audit

Prepared by:

- Control of Nonconforming Products
- Corrective Action
- Risk Management

Approved by:

Approved by:

DIRECTOR

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Excluded from the Scope of Quality Management System are the 2 Technical Services Divisions: The Standard and Testing Division (STD, the National Metrology Laboratory (NML), and the Advance Device Materials Testing Laboratory. Most of the laboratories of the STD are already certified under ISO 17025, starting with the Microbiology Laboratory in 2002 and followed by the Organic & Inorganic Chemistry Laboratories, and the Physical and Performance Testing Laboratory. Likewise, the National Metrology Laboratory is also ISO 17025 certified by Deutsche Akkreditierungsstelle (DAkks) of Germany since 2010.



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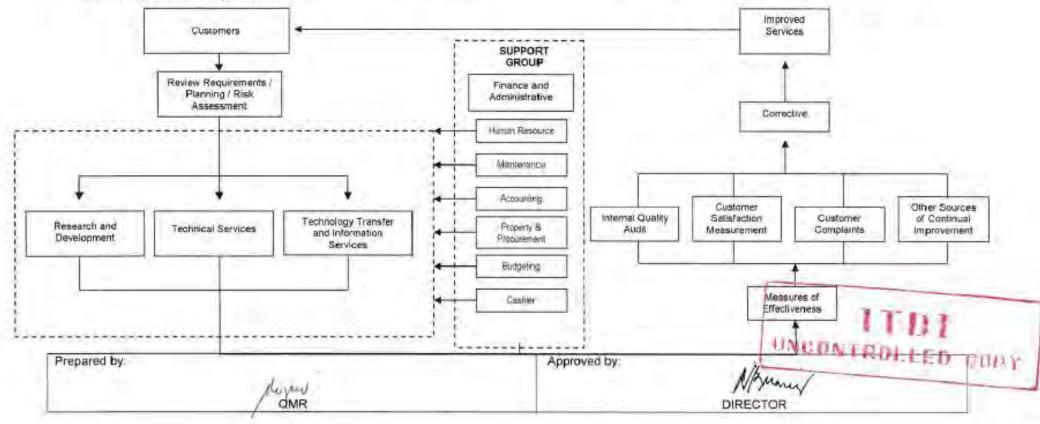
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SUBJECT	QUALITY MANAGEMENT SYSTEM AND ITS I	PROCESSES	

The QMS of ITDI covers processes that involve determining customer requirements, planning and managing the different activities involved to meet the requirements and expectations of its customers and conducting evaluation and improvement activities to enhance customer satisfaction. The diagram below illustrates the basic QMS processes.



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SECTION	LEADERSHIP	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	LEADERSHIP AND COM	MITMENT	

Top management has demonstrated leadership and commitment with respect to the quality management system by:

- Taking accountability for the effectiveness of the quality management system.
- Ensuring that the quality policy and quality objectives are established for the quality management system and are compatible with the context and strategic direction of the organization.
- Ensuring that integration of the quality management system requirements into the organization's business processes.
- · Promoting the use of the process approach and risk-based thinking.
- Ensuring that the resources needed for the quality management system are available.
- Communicating the importance of effective quality management and of conforming to the quality management system requirements.
- Ensuring that the quality management system achieves its intended results.
- Engaging, directing and supporting persons to contribute to the effectiveness of the quality management system.
- Promoting improvement.
- Supporting other relevant management roles to demonstrate their leadership as it applies to their areas of responsibility.

Customer Focus

Top Management has demonstrated leadership and commitment with respect to customer focus by ensuring that:

- Customer and applicable statutory and regulatory requirements are determined, understood and consistently met.
- The risks and opportunities that can affect conformity or products and services and the ability to enhance customer satisfaction are determined and addressed
- The focus on enhancing customer satisfaction is maintained.

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DIRECTOR

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DOCUMETED INFORMATION:

Quality Policy
Quality Objectives
Functional Objectives
PM-QMS 09-03 Management Review
Minutes of Management Review
PM-QMS 09-01 Customer Satisfaction Measurement



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SUBJECT	QUALITY POLICY		

Establishing the Quality Policy

Top management has established, implemented and maintained a quality policy that:

- Appropriate to the purpose and context of the organization and supports its strategic direction.
- Provides a framework for setting quality objectives.
- Includes a commitment to satisfy applicable requirements.
- Includes a commitment for continual improvements of the quality management system.

In pursuit of its commitment of providing customer satisfaction, Top Management has established a Quality Policy that reflects the values of employees of ITDI. The Quality Policy is stated as follows:

'We are committed to help local industries become globally competitive by providing appropriate technologies and services.

We shall continually improve our QMS to come up with conformity of products and services that would meet customer expectations within applicable regulatory and statutory requirements.

The quality objectives are:

- To complete and disseminate 90% of targeted R&D activities to meet the needs of local industries to become globally competitive.
- To deliver 90% of targeted technical services according to mutually agreed terms and conditions and existing applicable regulatory and statutory requirements.
- To attain at least a very satisfactory customer satisfaction rating for the year through the continual improvement of our QMS.

FUNCTIONAL OBJECTIVES:

Research & Development Group:

(Chemicals & Energy Division (CED) /

Environment & Biotechnology Division (EBD) /

Foods Processing Division (FPD) / Materials Science Division (MSD)

To implement 90% of approved projects for implementation based on agreed duration.

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| Director
| Director

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SUBJECT	QUALITY POLICY		

- To submit a terminal report two months after project completion.
- To implement 90% of the approved contract projects based on agreed schedule with Very Satisfactory customer rating.
- To undertake 90% of the approved technology transfer activities based on schedule with Very Satisfactory customer rating.

Research & Development Group: Packaging Technology Division (PTD)

- To implement on schedule 90% of the approved R&D projects and activities.
- To submit a terminal report two months after the project completion.
- To provide 90% of the request for packaging services with very satisfactory customer rating.

Support Group

Administrative Division - Cashier Section

 To ensure that 90% of approved disbursement vouchers are processed and transmitted to Agency's Government Servicing Bank within three (3) working days, subject to availability of funds

Administrative Division - Human Resource Management Section

- To ensure that 90% of requisitioned employees are hired within the prescribed period in accordance to Omnibus Rules and Regulations.
- To ensure that at least one (1) Learning and Development is attended by every staff within a
 year based on available resources.
- To ensure that 100% of IPCR submissions are monitored and reviewed in accordance with SPMS calendar and analyzed to identify developmental interventions.

Administrative Division - Property & Procurement Management Section

 To ensure that 90% of approved Purchased / Work Orders are delivered within the agreed delivery terms based on the specifications of the end-user and in accordance with the provisions of RA 9184.

 To ensure that 90% of newly acquired, donation and transferred properties are properly accounted, labeled and recorded within fifteen (15) days upon inspection and acceptance of the end-user.

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QMR	DIRECTOR

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SUBJECT	QUALITY POLICY		

To ensure that 100% of unserviceable properties are accounted and properly disposed within the scheduled plan in accordance with COA Rules and Regulations.

Administrative Division - Records Management Section

- To ensure that 90% of records received are properly disseminated within the day and shall be maintained and readily available when needed.
- To ensure that 90\% of records are controlled and disposed in accordance to provisions of RA 9470

Finance & Management Division

- To ensure that all ITDI activities with budgetary requirements are acted upon within the prescribed period.
- To ensure that all financial transaction are processed within the prescribed period.
- To ensure that all budgetary request and financial transactions conform with accounting and auditing rules and regulations within the prescribed period.

Planning and Management Information Systems Division

- To ensure the preparation of the institute's overall plans and programs for implementation during the year.
 - 90% of the Institute's programs and projects aligned to the national and department priorities and needs
 - 90% of the Institute's plans and budget proposals prepared within the prescribed date.
- To ensure Monitoring and Evaluation (M&E) of the Institute's programs, activities and projects (PAPs) during the year.
 - 2a Monitoring
 - 90% of projects monitored one month after the quarter.
 - 90% of projects monitored quarterly for risks identified.
 - 90% of quarterly reports prepared.
 - 2b. Evaluation
- UNCONTR 90% of project proposals evaluated with the ITDI-Technical Review Committee within the prescribed period.
 - 90% of ongoing projects evaluated with the ITDI-Technical Review Committee within two months after the quarter.

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2c. Agency Performance Review

- Preparation of the semestral Agency Performance Review Report two months after each semester with 100% accuracy.
- To ensure the review of existing and/or formulation of new organizational policies for improved operations and processes and effective governance with 100% acceptability.
- To ensure the implementation of appropriate information systems for the projects, services and other activities of the Institute as identified in the approved Information Systems Strategic Plan (ISSP, 2020-2022) and in accordance with the approved budget allocation; and
- To ensure action / response to requests for technical support on all ICT components:
 - 90% of requests for computer network technical support rendered within the agreed period.
 - 90% of requests for hardware technical support rendered with the agreed period.
 - 90% of requests for ICT System maintenance and support rendered within the agreed period.
 - 90% of requests for email technical support rendered within the agreed period.
 - 90% of requests for website maintenance and support rendered within the agreed period.

Technological Services Division

- To design and validate one new and acceptable training program within the year.
- To conduct 100% of the approved number of training programs for the year with a very satisfactory rating.
- To implement 100% of the approved requests for technical assistance / technology transfer in the Regions based on agreed schedules/duration within the year with a very satisfactory rating.
- To provide 100% relevant information to customers and stakeholders, both internally and externally in all forms (print and non-print, media / press conference, exhibits, techno fora) at the right time.
- To prepare techno-economic studies (market/industry profile, FS / BOP and techno-packages)
 of identified mature / commerciable technologies within the prescribed period.
- To develop business development strategies for marketing and promotion or identified technologies / new services within the prescribed period.

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/ QMR	DIRECTOR

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SUBJECT	QUALITY POLICY		

 To provide engineering related services in support of ITDI activities such as: R&D, technical services, maintenance and repair of physical facilities/buildings and technology transfer within the prescribed time with a very satisfactory rating.

Quality Management System:

Internal Quality Audit

- To ensure that internal audits shall be conducted once a year in accordance with ITDI's annual
 audit program. A detailed audit plan shall state the assignment of auditors in audit schedules as
 confirmed by auditees to determine whether the quality management system of ITDI:
 - Conforms to the planned arrangements of ISO 9001:2015; and
 - Is implemented and maintained.
- To make sure that the audit shall be carried out in accordance with 9.2 of ISO 9001;2015, taking
 into consideration the status and importance of the processes in the areas to be audited, and the
 results of the previous audits with corrections and corrective actions.
- To render audit reports stating conformance or non-conformance to ISO 9001 2015, and recommendations to areas for improvement as inputs for management review. A follow-up of corrective and preventive actions for nonconformities, monitor and review of internal audit program shall be done in accordance with PM-QMS 09-02.

Document Custodian

- To ensure that 100% of ITDI QMS documents are updated, maintained and controlled in accordance with its effectivity date.
- To ensure that 100% of ITDI QMS documents are properly issued on its effectivity date and withdrawn from the copy holder at the right time.

DOCUMENTED INFORMATION:

PM-QMS 09-03 Management Review Minutes of Meeting



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dopa	DIRECTOR
QMR	DIRECTOR

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SECTION	POLICY	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	ORGANIZATIONAL ROLES, RESPONSIBILITIES		ILITIES AND

Roles, Responsibilities and Authorities

Top management ensures that the responsibilities and authorities for relevant roles are assigned, communicated and understood within the organization through issuances of Office Orders, memorandums and during meetings in order to implement and maintain effective and efficient quality management system. The responsibilities and authorities of all employees throughout the organization are documented in the Manual of Actual Duties and Responsibilities. These are communicated to all employees during their orientation or their assignment to new positions and/or whenever there are changes to the position descriptions to enable them to contribute to the achievement of the quality objectives and to establish their involvement, motivation and commitment.

Top management assigns the responsibilities and authorities for

- Ensuring that the quality management system conforms to the requirements of the International Standard.
- Ensuring that the processes are delivering their intended outputs.
- Reporting on the performance of the quality management system and addressing opportunities for improvement, in particular to top management.
- Ensuring the promotion of customer focus throughout the organization.
- Ensuring that the integrity of the quality management system is maintained when changes to the quality management system are planned and implemented.

The Director designated the Chief Administrative Officer of ITDI to discharge the functions as the Quality Management Representative, on her behalf.

The responsibilities and authorities of the key personnel in the implementation and maintenance of the quality management system are described below:

Director

- Formulates the organizational plans and methods for the internal operations of ITDI;
- Chairs the ISO management review meetings; and
- Provides the necessary resources for the effective implementation and maintenance of the quality management system.

ISO Technical Committee

- Acts as the policy-making body responsible for setting the direction and overseeing the implementation of QMS activities;
- Reviews the effectiveness and suitability of the Institute's QMS;
- Approves QMS related final deliverables; and

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SUBJECT	ORGANIZATIONAL ROLES, RESPONSIBILITIES AND AUTHORITIES		ILITIES AND
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 Conducts ISO management reviews on such agenda as nonconformities, audit results, quality problems, performance indicators, corrective and preventive actions, and quality objectives.

Quality Management Representative (QMR):

Chief, Planning and Management Information Systems Division

Deputy QMR:

Chief, Technological Services Division

Members:

Chief, Chemicals and Energy Division

Chief, Environment and Biotechnology Division

Chief, Food Processing Division Chief, Materials Science Division

Chief, Finance and Management Division Chief, Packaging Technology Division

Chief, Administrative Division

Quality Management Representative

- Ensures that the quality management system is established, implemented and maintained in accordance with ISO 9001:2015 standard;
- Responsible for reporting and updating the top management on the performance of the quality management system for review, including recommendations for improvements;
- Ensures the promotion of awareness of customer requirements throughout the organization;
 and
- Acts as liaison with external parties on matters relating to quality management systems.

Division Chiefs

- Takes charge of the administrative operations of the division;
- Assist the QMR in ensuring that processes within the scope of the QMS are established, implemented & maintained;
- Report to the QMR on the performance of the QMS and any need for improvement;
- Assist the QMR in ensuring the promotion of awareness of customer requirements throughout the organization; and
- Ensure that all finished products conform to customer's specified requirements.

Section Chiefs

- Takes charge for the operations of the section;
- Reports to the Division Chief on the performance of the QMS and any need for improvement;
 and
- Assist the Division Chief in ensuring the promotion of awareness of customer requirements throughout the organization.

Decument Custodian Maintains and	controls QMS documents, including the re	vision, formal issuance, distribution
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Prepared by.	daya CMR	DIRECTOR DIRECTOR

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- Prepares Master Lists of Controlled Documents, Externally-Sourced Documents and Quality Records; and
- Serves as secretariat during Management Review and QMS related meetings.

Head Internal Audit Group

- · Prepares annual audit program
- · Ensures timely preparation of audit plan and audit checklist; and
- Implements the QMS internal audit based on the approved audit program.

Risk Review Committee

- Review and approve the existing divisional risk registers and the risk register crafted by the
 Risk Management Group
- Recommend changes, as needed, to the risk registers.
- Evaluate the effectiveness of the proposed changes to the risk registers.
- Monitor and evaluate the mitigating measures indicated by the Risk Management Group affecting the Institute's operational risks and strategies.
- Work with Risk Management Group in ensuring that the risks are properly considered and addressed

Human Resource Management Section

- Takes charge for recruitment, selection and placement of qualified personnel;
- Ensures the continuous supply of competent people to man the processes within the scope of the QMS.
- Ensures the effectiveness of the human resource interventions conducted for all personnel within the scope of the QMS;
- Ensures that relevant documents and records of employees are updated and filed;
- Responsible for the promotion of QMS in the institute;
- Identifies training needs of the Institute's employees in relation to their roles in the effective implementation and maintenance of the QMS; and
- Ensures that critical personnel and members are appropriately trained and competent to perform assigned task.

Property & Procurement Management Section

- Ensures that all purchases conform to the specified requirements and are acquired at the right time from the right source to the best advantage of the Center; and
- Ensures that suppliers are competent to provide quality products & services.

Technological Services Division

- Ensures that scheduled preventive maintenance of all equipment is implemented accordingly;
- Ensures the timely implementation of corrective maintenance.

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SUBJECT	ORGANIZATIONAL ROLES, RESPONSIBILITIES AND AUTHORITIES		

Budget and Cashier Section

Ensures that all finished job orders are properly billed and collected.

DOCUMETED INFORMATION:

Manual of Actual Duties and Responsibilities ITDI Special Order



Prepared by:	Approved by:
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/ QMR	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 06-01
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SECTION	PLANNING	PLANNING DOCUMENT TYPE	
SUBJECT	ACTIONS TO ADDR		AND

ITDI considered the issues referred to the organization and its context, the requirements referred to the needs and expectations of interested parties, and determine the risks and opportunities that need to be addressed to:

- Give assurance that the quality management system can achieve its intended results.
- Enhance desirable effects.
- · Prevent, or reduce, undesirable effects.
- Achieve improvement.

ITDI has planned:

a) Actions to address these risks and opportunities.

b) How to:

- Integrate and implement the actions into its quality management system processes.
- Evaluate the effectiveness of these actions.

Interested Parties	Requirements	Risks and Opportunities	Effectiveness of Actions
External Providers	~ Communication ~ Payment Terms ~ Timely delivery	~ Communication gap ~ Payment and delivery terms not specified ~ Delayed delivery	Communication like: email, phone, Payment and Delivery terms mention in PO
Customer	- Quality products and services - Timely delivery of product and services - Response to complaint Proper communication channel.	- Loss of trust - Loss of customer - Improve process and enhance customer satisfaction	 Defined in documented information and review in Management Review meetings.
Statutory and Regulatory Body	~ Complying with the statutory and regulatory requirements,	- output is not acceptable - aligned to government regulations, legal requirements, etc.	Defined in documented information and review in Management Review meetings.

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SUBJECT	ACTIONS TO ADDR		AND
SECTION	FEARITING	DOCUMENT TYPE	PAPER / PDF FILE
SECTION	PLANNING	EFFECTIVITY DATE	03 January 2020
	Quality Management System Manual	PAGE NUMBER	2 of 2
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Employees	~ Remuneration ~ Training and development	~ fair and just compensation and/or benefits. ~ Missed training opportunity due to limited budget	 compliance with the Magna Carta for S&T personnel. Defined in Staff development and review in Management Review meetings.
		~ technical expertise	1,

DOCUMENTED INFORMATION:

Risk Register Quality Objectives Functional Objectives Minutes of Meeting Minutes of Management Review



Prepared by:	Approved by:
Meigu	Manner
/ QMR	DIRECTOR

SUBJECT	QUALITY OBJECTIVES ACHIEVE		INING TO
SECTION		DOCUMENT TYPE	PAPER / PDF FILE
OFFICE	PLANNING	EFFECTIVITY DATE	03 January 2020
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ITDI has established quality objectives at relevant functions, levels and processes needed for the quality management system. The quality objective is stated as follows:

- To complete and disseminate 90% of targeted R&D activities to meet the need of local industries to become globally competitive.
- To delivery 90% of targeted technical services according to mutually agreed terms and conditions and existing applicable regulatory and statutory requirements.
- To attain at least a very satisfactory customer satisfaction rating for the year through the continual improvement of our QMS.

The quality objective is:

- · Consistent with the quality policy.
- Measurable.
- Take into account applicable requirements.
- Relevant to conformity of products and services and to enhancement of customer satisfaction.
- Monitored
- Communicated.
- Updated as appropriate.

In order to achieve its quality objectives, ITDI determined:

- What will be done?
- What resources will be required?
- Who will be responsible?
- When it will be completed
- How the results will be evaluated?



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/ QMR	DIRECTOR

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SECTION	PLANNING	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	QUALITY OBJECTIVES ACHIEVE		INING TO

DPCUMENTED INFORMATION:

Quality Objectives Functional Objectives Monitoring of Objectives



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SECTION	PLANNING	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	PLANNING OF	CHANGES	

ITDI has determined the processes needed for the quality management system and their application throughout the Institute's Quality Management System and its Processes.

When there are changes needed in the quality management system, ITDI determines such changes are carried out in a planned manner and considers the following:

- The purpose of the changes and their potential consequences.
- The integrity of the quality management system.
- The availability of resources.
- The allocation or reallocation of responsibilities and authorities.

DOCUMENTED INFORMATION:

QM-ITDI 04-04 Quality Management System and its Processes PM-ITDI 06-01 Change Management Process



Prepared by:	-	Reviewed / Approved by:
Ne	even)	Marine
/ (QMR .	DIRECTOR

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SUBJECT	RESOURCES		

General

ITDI determines and provides the resources needed for the establishment, implementation, maintenance and continual improvement of the quality management system. Thus, ITDI considers the capabilities of, and constraints on, existing internal resources and what needs to be obtained from external providers.

Top management ensures that the resources essential to the implementation of strategies and the achievement of the institute's objectives are identified and that the same are made available.

Decisions on the allocation of resources for operation and improvement of the QMS, enhancement of employees' competence through training, education and human resource intervention, the satisfaction of customers and other interested parties and planning for future resource needs are made during planning sessions. These resources may include people, infrastructure, work environment, information, suppliers and partners, and financial resources.

Plans are reviewed regularly to ensure the efficiency and the effectiveness of resource allocation.

People

ITDI ensures that personnel performing work affecting quality are competent on the basis of appropriate education, training and experience.

To assess, enhance and develop further their competence in providing quality products to the customers, the Administrative Division in coordination with the divisions concerned conducts annual training needs analysis based on competencies defined for each position. The Training Plan is prepared based on the identified training needs.

In addition to the provision of training, other actions and interventions are undertaken to address identified gaps in competence. Training and/or other actions taken are designed to ensure that all employees are aware of the relevance and importance of their activities and their contribution to the attainment of the quality objectives.

After each training or completion of other actions taken, the Chief, Administrative Division evaluates their effectiveness in improving competence of employees.

During the implementation of the QMS, all employees shall be appropriately provided with training and/or orientation to make them aware of the following:

- The importance of conformity with the quality policy and procedures and with the requirements of the quality management system;
- Their roles and responsibilities in achieving conformity with the requirements of the quality management system; and

Records of education, training, skills and experience of employees are included in the 201 file.



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SUBJECT	RESOURCES		

Infrastructure

ITOI determines, provides, and maintains the environment necessary for the operations of its processes and to achieve conformity of products and services. The infrastructure includes building and associated utilities, equipment, including hardware and software, transportation resources and information and communication technology.

Environment for the Operation of Processes

To achieve conformity of products and services, ITDI determines, provides and maintains a suitable environment which is combination of human and physical factors such as social, psychological, and physical environment.

ITDI also implements 5S program to provide employees opportunities for greater involvement thus, realizing the potential of people in the organization.

Monitoring and Measuring Processes

General

ITDI determines and provides the resources needed to ensure valid and reliable results when monitoring or measuring is used to verify the conformity of products and services to requirements. It ensures that the resources provided are suitable for the specific type of monitoring and measurement activities being undertaken and are maintained to ensure their continuing fitness for their purpose.

The institute retained documented information as evidence of fitness for purpose of the monitoring and measurement resources.

Measurement Traceability

To ensure valid results, measuring instruments and equipment are calibrated or verified or both at specified intervals and adjusted or re-adjusted as necessary. Measuring equipment is identified to determine the calibration status, safeguarded from adjustments that would invalidate the measurement result and be protected from damage and deterioration during handling, maintenance and storage.

In order to provide confidence in data, confirmation that the measuring devices are fit to use and are maintained to suitable accuracy and accepted standards, and identification of status are included in the measuring and monitoring processes.

When the equipment is found not to conform to requirements, concerned division/section assesses and records the validity of the previous measuring results.

Prepared by:

Approved by:

Approved by:

DIRECTOR

INI	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-01
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SUBJECT	RESOURCES		

Organizational Knowledge

The institute determines the knowledge necessary for the operation of its processes and to achieve conformity of products and services. This organizational knowledge based from internal and external sources are maintained and be made available to the extent necessary.

When addressing the needs and trends, ITDI considers its current knowledge and determines how to acquire or access any necessary additional knowledge and required updates, to enhance the effectiveness of the Quality Management System and enhance customer satisfaction.

DOCUMENTED INFORMATION:

Minutes of Planning Sessions Minutes of Staff Meetings Annual Training Plan PM-ADM-HRM 07-01 Human Resource Management PM-ADM-HRM 07-02 Staff Development PM-ADM-PPMS-08-01 Purchasing of Goods Training Record 201 File Preventive Maintenance Corrective Maintenance 5S Program Calibration Plan of Inspection, Measuring and Test Equipment Database of Intellectual Property Patents Certificates Terminal Reports Technical Papers Travel Reports Proceedings / Minutes of Planning Workshops



Prepared by	Approved by:
played	DIRECTOR

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		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	COMPETENCE	*	

ITDI has determined the necessary competence of person(s) doing work under its control that affects the performance and effectiveness of the quality management system. It ensured that these persons are competent on the basis of appropriate education, training or experience. Wherever applicable, the Institute takes action to acquire the necessary competence, and evaluate the effectiveness of the actions taken. These applicable actions can include the provision of training, mentoring or re-assignment of currently employed persons, or hiring or contracting of competent persons.

The Institute retained appropriated documented information as evidence of competence.

DOCUMENTED INFORMATION:

Training Record



Prepared by:	Reviewed / Approved by:
MR	DIRECTOR

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	SUPPORT	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	AWARENESS		

ITDI ensures that persons doing work under the Institute control are aware of the quality policy, the relevant quality objectives, their contribution to the effectiveness of the quality management system requirements and implications of not conforming with the quality management system requirements.

DOCUMENTED INFORMATION:

Quality Policy Quality Objectives Functional Objectives



Prepared by:	Approved by:
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/ QMR	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-04
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	SUPPORT	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	COMMUNICATION		

Pertinent information related to the effective implementation of the QMS are relayed to all employees through Office Orders, memoranda, meetings and posting at strategic places.

Internal communication to all levels and functions of the institute shall include but not limited to the following:

- Quality Policy:
- · Quality objectives:
- · Procedures relevant to their duties and responsibilities; and
- Results of monitoring, measurement, and analysis needed to ensure effective implementation.

Internal and external communications relevant to the quality management system are determined including; on what it will communicate, when to communicated, with whom to communicate, how to communicate and who communicates.

INTERNAL COMMUNICATION

WHAT	WHEN	WITH WHOM	HOW	MHO
Quality Policy / Quality Objectives	Permanent	All Employees Interested Parties	Display / Paper / Electronic	Director/QMR/
Effectiveness of QMS (QM/PM/WI)	Training / Orientation	All Employees	Paper / Electronic	Director / QMR DC
Internal Audit Plan	Before the Audit (at least once a year)	Top Management Process Owners of all functional areas	Paper / Electronic	Internal Quality Auditors
Management Review	After the Internal Audit	ITDI EXECOM	Meeting	Director
Strategic Plan - Approved Projects - Monitoring & Evaluation	First Quarter Succeeding month after each quarter	Division Heads / Project Leaders / Project Staff	Letter / email	PMISD
Roles, Responsibilities and Authorities	During recruitment / Promotion / Department change	Employee	Procedure / Oral / Training	Director ADM-HRMS
ITDI Office Order	When necessary	All Employees	Letter / Meeting	Director
Competency ~ Training needs ~ Training Plan	First quarter	ITDI Staff	Paper	Division Heads

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	SUPPORT	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	COMMUNICATION		

Customer Complaint / Feedback	When there are issues or rating is below VS.	Customer	Letter / Meeting	Process Owner / Division concerned / QMR / Director
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EXTERNAL COMMUNICATION

WHAT	WHEN	WITH WHOM	HOW	WHO
Information to External Providers	During pre- conference bidding / Submission of Quotation	External Providers	Purchase Order / Meeting / Telephone / Website / Email	Property & Procurement Management Section / Bids & Awards Committee / PMISD- Management Information System
Products and Services Information	During inquiry	Customer	MOA / Letter / Meeting	Division Chief / Project Leader / Staff
incoming and Outgoing Correspondences	When necessary	All Employees / Interested parties	Letter / email	Process Owner
Customer Feedback / Action Taken for Customer Complaints	When there are issues or rating is below VS.	Customer	Letter / Meeting	Process Owner / Division concerned / QMR / Director

DOCUMENTED INFORMATION:

Quality Policy
Quality Objectives
Functional Objectives
Office Orders
Incoming/Outgoing Correspondences
Minutes of Meeting
Records of Orientation/Re-orientation



Prepared by:	Approved by:
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QMR	DIRECTOR

	DEPARTMENT OF SCIENCE AND TECHNOLOGY INDUSTRIAL TECHNOLOGY DEVELOPMENT INSTITUTE	DOCUMENT CODE	QM-ITDI 07-05
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	REQUIREMENTS	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	DOCUMENTED INFORMATION		

Documentation of ITDI QMS is embodied in the documented statements of quality policy, objectives, quality manual, the mandatory processes required by ISO 9001:2015 and other documents needed by ITDI to ensure the effective planning, operation and control of the identified processes.

The Quality Management System Manual includes:

- The scope of the QMS as well as the details and justification for the exclusions in the QMS requirements;
- . The linkage of the procedures to the QMS; and
- The description of the interaction between the processes of the QMS.

The hierarchy of the QMS documentation is shown below:



Required maintain and retain documented information are maintained as evidence of implementation.

The QMR is responsible for the implementation of this policy.

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SECTION	QUALITY MANAGEMENT SYSTEM REQUIREMENTS	EFFECTIVITY DATE	03 January 2020
		DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	DOCUMENTED I	NFORMATIC	N

Creating and Updating

ITDI ensures that internally-generated and documents of external origin (in both hard copy and electronic form) needed for the effective implementations of its QMS are controlled. A documented procedure is maintained by the Document Custodian of the Institute to ensure appropriate:

- · Identification and description (e.g.a title, date, author, or reference number).
- Format (e.g. language, software version, graphics) and media (e.g. paper, electronic).
- Documents are reviewed and approved for suitability and adequacy.
- A Master List of Controlled Maintain Documented Information is maintained by DCI and DCD to prevent the use of obsolete documents. The list includes documents needed for knowledgepreservation and legal purposes. Such documents are properly marked.

Control of Documented Information

ITOI maintains and retains documented information to provide evidence of conformance to ISO 9001:2015 requirements and implementation of the QMS. All documented information required by the quality management system and by International Standard are controlled to ensure:

- It is available and suitable for use where and when it is needed.
- It is adequately protected (e.g. from loss of confidentiality, improper use, or loss of integrity).

For the control of documented information, ITDI is address the following:

- Distribution, access, retrieval and use.
- Storage and preservation, including preservation of legibility.
- Control of Changes (e.g. version control).
- Retention and disposition.

A Master List of Retain Documented Information is maintained to ensure control. Maintenance and storage of confidential records are the responsibility of the concerned heads.

Documented information of external origin determined by the organization to be necessary for the planning and operation of the quality management system is identified as appropriate, and be controlled.

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			ITDI	DIRECTOR

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SUBJECT	DOCUMENTED INFORMATION		N

Documented information retained as evidence of conformity is protected from unintended alterations.

DOCUMENTED INFORMATION:

PM-QMS 07-01 Control of Maintain Documented Information
PM-QMS 07-02 Control of Retain Documented Information
PM-QMS 07-03 Control of Maintain and Retain Documented Information in Electronic Media
Master List of Controlled Maintain Documented Information
Master List of Quality Retain Documented Information
Master List of Externally-Sourced Documented Information



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/ QMR	DIRECTOR

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	OPERATION	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	OPERATIONAL PLANNING AND CONTROL		

ITDI has planned, implemented and control the processes needed to meet the requirements for the provision of products and services, and implemented the actions determined in planning requirements for the products and services are determined, established criteria for the processes and acceptance of products and services; determined the resources needed to achieve conformity to the product and service requirement; implemented control of the processes in accordance with the criteria; and determined, maintained and retained documented information to the extent necessary to have confidence that the processes have been carried out as planned and demonstrated the conformity of products and services to the customer's requirements.

The output of this planning is suitable for the Institute's operations.

ITDI has controlled in changes of the plan, reviewed the consequences of unintended changes, take action to mitigate any adverse effects, as necessary. It ensure that outsourced processes are controlled.

DOCUMENTED INFORMATION:

PM-ITDI 08-01 Provision of Technical Services

PM-ITDI 08-02 Generation and Approval of Regular R&D Projects

PM-ITDI 08-03 Implementation of Approved Regular R&D Projects

PM-CED 08-01 Energy Audit

PM-EBD-CPS 08-01 Cleaner Production Assessment

PM-EBD-CPS 08-02 Environmental Technology Verification

PM-EBD-CPS 08-03 Rapid Cleaner Production Assessment

PM-PTD 08-01 Label Design Development

PM-PMISD 08-01 Planning, Monitoring and Evaluation of R&D Project

PM-TSD-BDS 08-01 Technology Transfer / Commercialization of ITDI Matured Technologies

PM-TSD-IDS 08-01 Creation and Production of Social Marketing Collaterals

PM-TSD-RCTS 08-01 Facilitating Regional Cooperation and Training

PM-TSD-RCTS 08-02 Planning and Scheduling of Regular / Funded Technology Transfer Program

PM-TSD-RCTS 08-03 Implementation of Technical and Other Regional Intervention

PM-TSD-RCTS 08-04 Planning and Scheduling of Regular Training Program

PM-TSD-RCTS 08-05 (implementation of Training)

PM-TSD-RCTS 08-06 Design and Development of New Training Program

PM-ADM-PPMS 08-01 Purchasing of Goods

PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased Items

PM-FMD-AS 08-01 Billing

PM-FMD-BS 08-01 Budget Preparation

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Prepared by:

Approved by:

Approved by:

DIRECTOR

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SUBJECT	REQUIREMENTS FOR PRO	DUCTS AND	SERVICES

Customer Communication

Communication with customers include:

- Providing information relating to products and services.
- Handling enquiries, contracts or orders, including changes.
- Obtaining customer feedback relating to products and services, including customer complaints.
- Handling or controlling customer property.
- Establishing specific requirements for contingency actions, when relevant.

Determining the Requirements for Products and Services

When determining the requirements for the products and services to be offered to customer, the institute ensures that:

- The requirements for the products and services are defined including any applicable statutory and regulatory requirements and those considered necessary by the organization.
- The Institute can meet the claims for the products and services it offers.

Review of the Requirements for Products and Services

iTDI ensures that it has the ability to meet the requirements for products and services to be offered to customers. It conducts a review before committing to supply products and services to a customer, to include:

- Requirements specified by the customer, including the requirements for delivery and postdelivery activities.
- Requirements not stated by the customer, but necessary for the specified or intended use, when known.
- Requirements specified by the Institute.
- Statutory and regulatory requirements applicable to the products and services.
- Contract or order requirements differing from those previously expresses.

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SUBJECT	REQUIREMENTS FOR PRO	DUCTS AND	SERVICES

ITDI ensures that contract or order requirements differing from those previously defined are resolved. The customer's requirements are confirmed by ITDI before acceptance, when the customer does not provide a documented statement of their requirements.

The Institute retains documented information, as applicable:

- · On the results of the review
- · On any new requirements for the products and services

Changes to Requirements for Products and Services

(TD) ensures that relevant documented information is amended, and that relevant persons are made aware of the changed requirements, when the requirements for products and services are changed.

DOCUMENTED INFORMATION:

PM-ITDI 08-01 Provision of Technical Services

PM-ITDI 08-02 Generation and Approval of Regular R&D Projects

PM-ITDI 08-03 Implementation of Approved Regular R&D Projects

PM-CED 08-01 Energy Audit

PM-EBD-CPS 08-01 Cleaner Production Assessment

PM-EBD-CPS 08-02 Environmental Technology Verification

PM-EBD-CPS 08-03 Rapid Cleaner Production Assessment

PM-PTD 08-01 Label Design Development

PM-PMISD 08-01 Planning, Monitoring and Evaluation of R&D Project

PM-TSD-BDS 08-01 Technology Transfer / Commercialization of ITDI Matured Technologies

PM-TSD-IDS 08-01 Creation and Production of Social Marketing Collaterals

PM-TSD-RCTS 08-01 Facilitating Regional Cooperation and Training

PM-TSD-RCTS 08-02 Planning and Scheduling of Regular / Funded Technology Transfer Program

PM-TSD-RCTS 08-03 Implementation of Technical and Other Regional Intervention

PM-TSD-RCTS 08-04 Planning and Scheduling of Regular Training Program

PM-TSD-RCTS 08-05 Implementation of Training

PM-TSD-RCTS 08-06 Design and Development of New Training Program

PM-ADM-PPMS 08-01 Purchasing of Goods

PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased items

PM-FMD-AS 08-01 Billing

PM-FMD-BS 08-01 Budget Preparation

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Prepared by:	Approved by:
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QMR	DIRECTOR

SUBJECT	DESIGN AND DEV PRODUCTS AN		
SECTION	OF ENATION	DOCUMENT TYPE	PAPER / PDF FILE
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General

ITDI has established, implemented and maintained a design and development process that is appropriate to ensure the subsequent provision of products and services.

Design and Development Planning

Top management ensures that the Research and Development Divisions such as CED, EBD, FPD, and MSD, PTD and concerned section of TSD of ITDI has determined the stages and controls for design and development.

For every R&D project and packaging design, the Project Team considers the following:

- The nature, duration and complexity of the design and development activities.
- The required process stages, including applicable design and development reviews.
- The required design and development verification and validation activities.
- The responsibilities and authorities involved in the design and development process.
- The internal and external resources needs for the design and development of products and services.
- The need to control interfaces between persons involved in the design and development process.
- The need for involvement of customers and user in the design and development process.
- The requirement for subsequent provision of products and services.
- The level of control expected for the design and development process by customers and other relevant interested parties.
- The documented information needed to demonstrate that design and development requirements have been met.

Whenever a customer requires a new technical training program that is not included in the master list of training program, the head of TSD and EBD/Design Team prepares a design and development plan. The plan describes the different activities and defines responsibility for their implementation. It also describes the training objectives and expected outputs, sequencing of the design and development process, design methodologies, review, verification and validation activities. The plan is updated whenever changes in the design occur.

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	OPERATION	DOCUMENT TYPE	PAPER / PDF FILE
SUBJECT	DESIGN AND DEVELOPMENT OF PRODUCTS AND SERVICES		

Design and Development Inputs

inputs to design and development include functional and performance requirements, information derived from previous similar design and development activities, statutory and regulatory requirements, standards or codes of practice that the Institute has committed to implement, potential consequences of failure due to the nature of the products and services.

Inputs is adequate for design and development purposes, complete and unambiguous. Conflicting design and development inputs is resolved.

Concerned sections of CED, EBD, FPD, MSD, PTD, and TSD ensure that all inputs are reviewed for adequacy and that requirements are clear, unambiguous and not in conflict with each other.

The Institute is retained documented information on design and development inputs.

Design and Development Controls

The Institute applies controls to the design and development process to ensure that:

- The results to be achieved are defined.
- Reviews are conducted to evaluate the ability of the results of design and development to meet requirements.
- Verification activities are conducted to ensure that the design and development outputs meet the input requirements.
- Validation activities are conducted to ensure that the resulting products and services meet the requirement
- Any necessary actions are taken on problems determined during the review, or verification and validation activities.
- Documented information of these activities is retained.

Design and Development Outputs

CED, EBD, FPD, MSD, PTD, and TSD Design Team ensure that output of design and development outputs meet the input requirements; adequate for the subsequence processes for the provision of products and services; include or reference monitoring and measuring requirements, as appropriate, and acceptance criteria; and specify the characteristics of the products and services that are essential for their intended purpose and their safe and proper provision.

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The Institute has retained documented information on design and development outputs. Design output includes data demonstrating the comparison of process inputs to process outputs, product specifications including acceptance criteria, process, material and testing specifications, training requirements, user information, purchase requirements, and qualification tests and reports.

Design outputs are reviewed against inputs to provide objective evidence that outputs have effectively and efficiently met requirements for the process and product,

Design and Development Changes

Design and development changes are identified, reviewed and control of changes made during, or subsequent to, the design and development of products and services, to the extent necessary to ensure that there is no adverse impact on conformity to requirements.

Documented information retained on: design and development changes, results of reviews, authorization of the changes, and actions taken to prevent adverse impacts.

DOCUMENTED INFORMATION:

Design and Development Plan
Design and Development Inputs
Design and Development Controls
Design and Development Changes
Results of Reviews
Authorization of Changes
Actions Taken to Prevent Adverse Impacts



Prepared by:	Approved by:
OMR	DIRECTOR

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SUBJECT	PROCESSES, PRODUC		

General

To ensure that externally provided processes, products and services conform to requirement, the Property & Procurement Management Section has determined the controls to be applied to externally provided processes, products and services when:

- Products and services from external providers are intended for incorporation into the Institute's own products and services.
- Products and services are provided directly to the customer(s) by external providers on behalf
 of the organization.
- A process, or part of a process, is provided by an external provider as a result of a decision by the Institute.

The Institute determines and applies criteria for the evaluation, selection, monitoring of performance, and re-evaluation of external providers, based on their ability to provide processes or products and services in accordance with requirements

The Master List of Accredited External Providers is retained documented information and their performance is evaluated annually through the use of rating sheet.

Whenever purchase requests are made, complete specification and description of items are indicated to ensure that only the required items are purchased. No purchases are made without the approval of the Director or her Authorized Representative. Personnel implementing the purchase are competent based on their education, training and experience.

In case of subcontracted services for some processes that the organization's capability may not be adequate, said services should be in conformance to the requirements of the customer.

Type and Extent of Control

To ensure that externally provided processes, products and services do not adversely affect the Institute's ability to consistently deliver conforming products and services to its customers, the Institute is:

- Ensure that externally provided processes remain within the control of its quality management system:
- Define both the controls that it intends to apply to an external provider and those it intends to apply to the resulting output.
- Take into consideration: the potential impact of the externally provided processes, products and services on the Institute's ability to consistently meet customer and

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SUBJECT	PROCESSES, PRODUC		

applicable statutory and regulatory requirements and the effectiveness of the controls applied by the external provider.

Determine the verification, or other activities, necessary to ensure that the externally
provided processes, products and services meet requirements.

Information of External Providers

To ensure the adequacy of requirement prior to the communication to the external providers, the Institute communicates its requirement for:

- The processes, products and services to be provided.
- The approval of products and services, methods, processes and equipment, and the release
 of products and services.
- Competence, including any required qualification of persons.
- The external providers' interactions with the organization
- Control and monitoring of the external providers' performance to be applied by the institute.
- Verification or validation activities that the organization, or its customer, intends to perform at the external providers' premises.

DOCUMENTED INFORMATION:

PM-ADM-PPMS 08-01 Purchasing of Goods PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased Items Master List of Accredited External Providers



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OMR	DIRECTOR

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SUBJECT	PRODUCTION AND SERVICE PROVISION		

Control of Production and Service Provision

ITDI implements production and service provision under controlled conditions that include, as applicable:

- The availability of documented information that defines: (1) the characteristics of the products
 to be produced, the services to be provided, or the activities to be performed; and (2) the results
 to be achieved.
- The availability and use of suitable monitoring and measuring resources.
- The implementation of monitoring and measurements activities at appropriate stages to verify that criteria for control of processes or outputs, and acceptance criteria for products and services, have been met
- The use of suitable infrastructure and environment for the operation of processes.
- The appointment of competent persons, including any required qualification.
- The validation and periodic revalidation, of the ability to achieve planned results of the
 processes for production and service provision, where the resulting output cannot be verified
 by subsequent monitoring or measurement.
- The implementation of actions to prevent human error.
- The implementation of release, delivery and post-delivery activities.

Identification and Traceability

The Institute uses suitable means to identify outputs when it is necessary to ensure the conformity of products and services.

It identifies the status of outputs with respect to monitoring and measurement requirements throughout production and service provision.

Provision of services is uniquely identified using Reference Number. The assigned Reference No. can be traced from the time the customer inquiry is received up to the time the product is released. The Reference/Project Number No. is indicated in all records.

ITDI has retained the documented information necessary to enable traceability:

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SUBJECT	PRODUCTION AND SERVICE PROVISION		

Property Belonging to Customers or External Providers

ITDI exercises care with materials, components, tools and equipment, premises, intellectual property and personal data belonging to customer or external providers while it is under the institute's control or being used by the institute. It is identified, verified, protected and safeguarded. If any customer property is lost, damaged or found to be unsuitable for use, the customer is informed and retained documented information on what has occurred.

Preservation

Conformity of product during production and service provision and release to the intended destination are preserved by the ITDI in order to maintain conformity to requirements. Appropriate labels, handling instructions, contamination control, packaging, storage, transmission or transportation, and protection are ensured and protected to prevent damage during transport.

Post - Delivery Activities

To meet requirements for post-delivery activities associated with the products and services. ITDI determines the extent of post-delivery activities that are required and considers:

- Statutory and regulatory requirements.
- The potential undesired consequences associated with its products and services.
- The nature, use and intended lifetime of its products and services.
- Customer requirements.
- Customer feedback

Control of Changes

To ensure continuing conformity with requirements, the Institute reviews and controls changes for production or service provision, to the extent necessary to ensure continuing conformity with requirements.

Documented information describing the results of the review changes, the person(s) authorizing the change, and any necessary actions from the review are retained.

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DOCUMENTED INFORMATION:

PM-ITDI 08-01 Provision of Technical Services

PM-ITDI 08-02 Generation and Approval of Regular R&D Projects

PM-ITDI 08-03 Implementation of Approved Regular R&D Projects

PM-CED 08-01 Energy Audit

PM-EBD-CPS 08-01 Cleaner Production Assessment

PM-EBD-CPS 08-02 Environmental Technology Verification

PM-EBD-CPS 08-03 Rapid Cleaner Production Assessment

PM-PTD 08-01 Label Design Development

PM-PMISD 08-01 Planning, Monitoring and Evaluation of R&D Project

PM-TSD-BDS 08-01 Technology Transfer / Commercialization of ITDI Matured Technologies

PM-TSD-IDS 08-01 Creation and Production of Social Marketing Collaterals

PM-TSD-RCTS 08-01 Facilitating Regional Cooperation and Training

PM-TSD-RCTS 08-02 Planning and Scheduling of Regular / Funded Technology Transfer Program

PM-TSD-RCTS 08-03 Implementation of Technical and Other Regional Intervention

PM-TSD-RCTS 08-04 Planning and Scheduling of Regular Training Program

PM-TSD-RCTS 08-05 Implementation of Training

PM-TSD-RCTS 08-08 Design and Development of New Training Program

PM-ADM-PPMS 08-01 Purchasing of Goods

PM-ADM-PPMS 08-02 Acceptance, Storage and Release of Purchased Items

PM-FMD-AS 08-01 Billing

PM-FMD-BS 08-01 Budget Preparation

PM-QMS 08-01 Handling Customer Complaint

PM-QMS 08-02 Control and Monitoring of Customer Supplied Materials



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SUBJECT	RELEASE OF PRODUC	TS AND SE	RVICES

ITDI implements planned arrangements, at appropriate stages, to verify that the product and service requirements have been met.

The release of products and services to the customer shall not proceed until the planned arrangements have been satisfactorily completed, unless otherwise approved by a relevant authority and, as applicable, by the customer.

The Institute retained documented information on the release of products and services that include:

- Evidence of conformity with the acceptable criteria
- Traceability to the person(s) authorizing the release.

DOCUMENTED INFORMATION:

Records of Traceability / Logbook



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SECTION	OPERATION	EFFECTIVITY DATE	03 January 2020
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SUBJECT	CONTROL OF NONCONE	ORMING OF	UTPUTS

ITDI ensures that product which does not conform to product requirements is identified and controlled to prevent its unintended use or delivery. It takes appropriate action based on the nature of the nanconformity and its effect on the conformity of products and services. It also applies to nonconforming products and services detected after delivery of products, during or after the provision of services.

The Institute deals with nonconforming outputs in one or more of the following ways:

- Correction
- Segregation, containment, return or suspension of provision of products and services.
- Informing the customer.
- Obtaining authorization for acceptance under concession.

Top management empowers the employees with the authority and responsibility to report nonconformities at any stage of a process in order to ensure timely detection and disposition of nonconformities. Authority for response to nonconformities and actions to be taken are defined in the documented procedures to maintain achievement of process and product requirements.

Use of, release and acceptance of nonconforming products is done under concession by divisions/sections chief concerned and, where applicable, by the customer.

All chiefs of divisions and sections concerned ensure that appropriate action be taken to preclude the original intended use or application of any nonconforming product.

Whenever applicable, all chiefs of divisions and sections concerned ensure that appropriate action be taken to the effects or potential effects of the non-conformity when non-conforming product is detected after delivery or use has started.

Conformity to the requirements is verified when nonconforming outputs are corrected.

The Institute retained documented information that: described the nonconformity, describes the actions taken, describes any concessions obtained, and identifies the authority deciding the action in respect of the nonconformity.

DOCUMENTED INFORMATION:

PM-QMS 09-03 Control of Nonconforming Outputs



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SUBJECT	MONITORING, MEASURE EVALUA		LYSIS AND

General

Top management ensures the effective and efficient measurement, collection and validation of data to ensure conformity to product requirements as well as the customers' level of satisfaction. The validity and purpose of measurements and the intended use of data are reviewed to ensure that these processes add value to the Institute.

Performance of the Institute's processes include measurement and evaluation of its services, capability of its processes, achievement of objectives, targets and programs, compliance to applicable legal and other requirements and satisfaction of customer.

Data derived from the monitoring of performance provide information and serves as guide in its continual improvement activities. Results of the analysis of data from improvement activities is considered as one input to management review in order to provide information for improving the performance of the Institute.

Applicable methods including statistical techniques and the extent of their use are included.

ITDI is evaluating the performance and the effectiveness of the quality management system. The QMR in consultation with all the respective divisional chiefs, implements the monitoring, measurement, analysis and improvement processes needed, as to:

- What needs to be monitored and measured.
- The method for monitoring, measurement, analysis and evaluation needed to ensure valid results.
- When the monitoring and measuring shall be performed.
- When the results from monitoring and measurement are analyses and evaluated.

The institute is retaining appropriate documented information as evidence of the results.

Customer Satisfaction

In order to measure the performance of the QMS, information relating to customer perception of the degree to which their needs and expectations have been fulfilled is monitored through the conduct of Customer Satisfaction Survey. The Institute determines the methods for obtaining, monitoring and reviewing this information.

The QMR in coordination with the Division Chiefs are responsible for Customer Satisfaction as one of the measurements of the Quality Management System and as to whether the Institute has mel customer requirements. Customer satisfaction shall include collection of authentic data, frequency and validity of analysis. The trend and major elements affecting customer satisfaction and key indicators of customer dissatisfaction will be monitored supported by objective evidence.

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A customer satisfaction survey is done periodically and its analysis and presentation is done annually and presented in the Management Review. Any negative feedbacks are addressed immediately.

Analysis and Evaluation

ITDI analyzes and evaluates appropriate deta and information arising from monitoring and measurement. The results of analysis are used to evaluate:

- Conformity of products and services
- The degree of customer satisfaction.
- The performance and effectiveness of the quality management system.
- If planning has been implemented effectively.
- The effectiveness of action taken to address risks and opportunities
- The performance of external providers.
- The need for improvements to the quality management system.

Monitoring and Measurement of Processes

Suitable methods for monitoring and, where applicable, measurement of the quality management system processes are applied to demonstrate the ability of the processes to achieve planned results. When planned results are not achieved, appropriate correction and corrective action are taken to ensure conformity of the product.

Monitoring and Measurement of Product

To verify that product requirements have been met, the Institute ensures that monitoring and measurement are carried out at appropriate stages of the product realization process in accordance with the planned arrangements.

Evidence of conformity with the acceptance criteria is maintained. Records indicate the person(s) authorizing the release of product. Only products which have been satisfactorily completed through the planned arrangements are released to the customer

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DOCUMENTED INFORMATION:

Accomplished Customer Feedback Survey Form Analysis of Data File/Report Plan for Preventive Action Action Plan

Monitoring of Quality Objectives



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SUBJECT	INTERNAL AUDIT		

Internal Audit

The Internal Audit Committee is responsible in planning and implementing internal audits in order to verify whether the quality management system conforms to the planned arrangements, to the requirements of ISO 9001:2015 and to the quality management system requirements established by the Institute and whether the QMS is effectively implemented and maintained.

An audit program is planned and conducted at least once a year, taking into consideration the status and importance of the processes and areas to be audited, as well as the results of previous audits. The Audit Plan defines the audit criteria, scope, frequency and methods of audit. Auditors are selected based on competence, training, work experience and qualification and shall not audit their own work. Objectivity and impartiality of the audit process is ensured in the selection of auditors and conduct of audits.

The responsibilities and requirements for planning and conducting audits, and for reporting results and maintaining records are defined in PM-QMS 09-02, Internal Audit

The Heads responsible for area being audited ensure that any necessary corrections and corrective actions are taken without undue delay to eliminate detected non-conformities and their causes.

Follow-up activities include the verification of the actions taken and the reporting of the verification results. Internal audit reports include evidence of excellent performance in order to provide opportunities for recognition by management and motivation of people.

The Institute has retained documented information as evidence of the implementation of the audit programme and the audit results.

DOCUMENTED INFORMATION:

PM-QMS 09-02 Internal Audit Audit Program Audit Results



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MANAGEMENT REVIEW		
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Top Management, together with the Quality Management Representative, concerned divisions and section chiefs reviews the effectiveness of the QMS at least once a year to ensure its continuing suitability, adequacy, effectiveness and alignment with the strategic direction of the organization. The review involves identifying opportunities for improvement and possible need for changes, including the quality policy and objectives.

The management review is planned and carned out taking into consideration of inputs:

- a) The status of actions from previous management review.
- b) Changes in external an internal issues that are relevant to the quality management system.
- c) Information on the performance and effectiveness of the quality management system, including trends in:
 - Customer satisfaction and feedback from relevant interested parties.
 - 2. The extent to which quality objectives have been met.
 - 3. Process performance and conformity of products and services.
 - Nonconformities and corrective actions.
 - Monitoring and measurement results.
 - 6. Audit results.
 - 7. The performance of external providers
- d) The adequacy of resources.
- e) The effectiveness of actions taken to address risks and opportunities.
- f) Opportunities for improvement.

Outputs of the management review include opportunities for improvement, any need for changes to the quality management system and resource needs.

Retain documented information from management reviews are maintained to provide for traceability and to facilitate evaluation of the management review process itself, to ensure its continued effectiveness and added value to the organization.

DOCUMENTED INFORMATION:

PM-QMS 09-01 Management Review

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SUBJECT	IMPROVEMENT		

General

ITDI determines and selects opportunities for improvement and implements any necessary actions to meet customer requirements and enhance customer satisfaction which include:

- Improve products and services to meet requirements as well as to address future needs and expectations.
- · Correct, prevent or reduce undesired effects.
- Improve the performance and effectiveness of the quality management system.

Non Conformity and Corrective Action

When a nonconformity occurs, including any arising from complaints, the QMR ensures that all nonconformities relating to product, process and quality system are investigated and the results are recorded. While investigating causes of non-conformances and deciding corrective actions, the QMR identifies other products / situations where same or similar nonconformance can occur.

Corrective actions for nonconforming products and processes, detected in the organization are recorded, analyzed, reviewed, and necessary actions are taken. This will be the responsibility of process owner. The corrective action and their effectiveness is monitored.

Documented procedures are established to define requirements for corrective action and for handling customer complaints.

Corrective actions are initiated whenever there are complaints from customers; audit findings; performance standards, objectives and targets are not met within the defined time frame; and nonconforming products, both internal and external to the organization occur.

Causes of nonconformities are analyzed and took suitable action to prevent their reoccurrence. The effectiveness of the identified corrective action is monitored through customer feedback and also the customers are kept aware of the corrective actions initiated, with each identified corrective action.

Continual Improvement

Data generated from the monitoring and measurements of processes are analyzed regularly to determine areas that need to be continually improved to ensure effective implementation of QMS. This information is taken from the regular reviews of the quality policy, objectives, audit results, corrective and preventive actions, results of management reviews and analysis of customer feedback. The concerned heads are responsible for ensuring that continual improvement activities are carried out.

The documented procedure has defined requirements for the determination of potential nonconformities and their causes, preparation and implementation of preventive actions where



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needed, and review effectiveness of actions taken. The Division Chiefs and QMR maintain records of actions taken and is presented during management review meetings.

DOCUMENTED INFORMATION:

PM-QMS 08-01 Handling Customer Complaint PM-QMS 09-04 Corrective Action



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QMR	DIRECTOR