Seminar on

Scheme for approval of IT test Laboratories

for

eGovernance Solutions

Welcome to the Participants

17.01.2014 Bangaluru



STQC

Standardisation

Testing

Quality

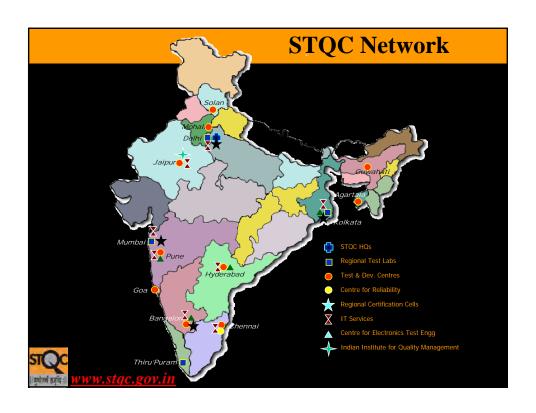
Certification



Standardisation Testing & Quality Certification Directorate
An attached office of DeitY



www.stac.gov.ii



Our - Objective

STQC was setup under Department of Electronics as a promotional programme 1977. It was upgraded to a level of attached office of DOE 1985 with the following objective :

" To assist small and medium enterprises in electrotechnical sector in improving the quality of their products through Testing & Calibration services."

In tune with the current needs when DOE became Department of Electronics & Information Technology , STQC has redefined its objective:

"To be a key enabler in making Indian organisations and IT users achieve compliance to International Quality Standards and compete globally"

STQC II minimi anik:

www.stqc.gov.in

STQC - Shifting Focus

- STQC is shifting its Focus from Service Provider to Framework Provider
- This will provide opportunities for private Test Houses to Network with STQC and explore new business opportunities
- STQC will focus on Certification and use empanelled test laboratories as its resource centres
- Capacity Building will be priority area for STQC

Following Training Programme in this area will be conducted:

- ➤ Overview of ISO/IEC 17025 for S/W Test Lab
- ➤ Implementation of ISO/IEC 17025 for S/W Test Lab
- ➤ Assessor Programme for ISO/IEC 17025 in association with NABL



Approval and Accreditation

- STQC has designed an approval scheme for independent IT Test Labs
- The scope is Testing of eGovernance Solutions
- STQC intends to recognize and empanel private test houses by means of approval scheme for Testing eGovernance Software
- STQC is not an accreditation Body and does not want to have conflict with NABL which is National Accreditation Body for Laboratories and work under DST
- This Software Test House approval Scheme is designed to meet the objective of DeitY and to have Quality assessed Software for eGovernance



Agenda Conformity Assessment in eGovernance - U K Nandwani Introduction to the Approval Scheme for ITTL - U K Nandwani Requirements of ISO/IEC 17025 for Software Lab - B. Singh Experience sharing in IT Test Laboratory accreditation - B. S. Kumar Panel Discussion

Contact us

My team can be contacted for any query in this regard and in case required I am available at –

email: dgstqc@deity.gov.in



Conformity Assessment in eGovernance

17.01.2014 Bangaluru

eGovernance

eGovernance is transformation of internal and external relationship of Government Sector, by using information and communication technologies. This is by re-engineering government service delivery process and aligning it with IT leading to increase an efficiency, effectiveness and transparency.

e-Governance - Need for Conformity Assessment

Benefits

Reduce service time

Improved customer service through up-to-date, accurate data.

Business intelligence for fact based decision making

Increased Government revenue due to reduction in transmission and distribution losses

Risk

Economic Risk

- Huge Investment
- Cost of Technology and Knowledge is high

Technological Risk

- High obsolesce Rate
- Dependability/Reliability of Technology
- Use of right technology

Social Risk and User acceptability Risks

- Solutions are citizen & business Centric & touch sensitive service oriented issues
- High expectation

Concerns

• Users

- whether Government services will be available in a convenient way as promised

· Policy Makers and Administrators

-Whether objectives of eGovernance are being achieved (Transparency, availability of Service, compliance with Govt. Rules, procedures, decisions and Regulations)

• Solution/Service Provider

-That system meets the requirements of RFP.

eGovernance Conformity Assessment-Business of Confidence

To have confidence on the eGovernance solution,
Administrators and Funding agencies look for an affirmative indication or judgment by independent agency that a product or service has met the requirements of a contract and applicable regulation. The process for determining the degree of compliance of the solution characteristics (as delivered) with the requirements (as desired) by means of objective evaluation is known as a the conformity assessment.

It provides the information on the fulfillment of the provisions of "Request for Proposal" and "Contract Agreement". The techniques used for Conformity Assessment are sampling, testing, inspection, review, assessment, evaluation and certification.

Conformity Assessment - Purpose

Business of Confidence

To Administrator

Achieving his vision

Addressing his concern

Strategies handling of complex problems

To solution providers

A well accepted approach based on ISO standards.

Common methodology and criteria - fair playground for all solution providers

Independent third party

To Users

Solution is usable, secure and meet Service Level Agreements Solution is functional and legally compliant

Benefits and Risks

When applied correctly, conformity assessment can

- Provide purchaser, a confidence in the suppliers, whose services they purchase
- Help to assess the strengths and weaknesses of the initiatives and projects
- Reduce Risks by providing inputs for early monitoring and timely corrections
- Help funding agencies to get confidence on the outcome of the project
- Help businesses to be competitive
- Facilitate trust in procurement and supply
- Provide a visible link between standards and the market

However, if applied incorrectly, conformity assessment can also...

- Be a burden on business by adding cost of demonstrating compliance
- Create barriers to procurement and supply
- Inhibit innovation
- Confuse the market

Conformity Assessment

e GCA - Objective

Generating Confidence of Citizen and Business on e-Government

Through conformity assessment to user-requirements, regulations and Best Practices by Independent Third Party

Rather than

Relying solely on the assertion of the developers and solution providers

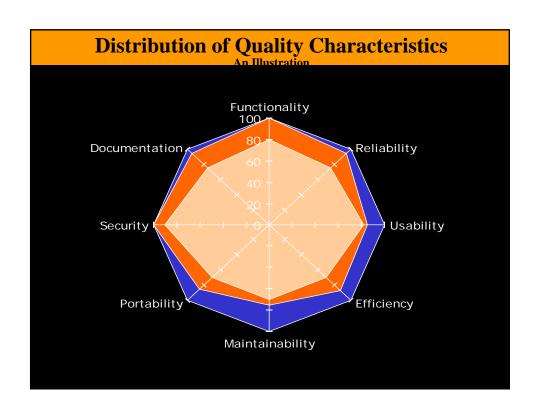
eGCA- Quality Gates

eGovernance Conformity Assessment has four Quality Gates

- Software Application Quality
- System and Infrastructure
- Service Quality (SLA)
- Quality Processes in Government Organisation

Quality Gate 1 – Software Quality

- Application Quality is basic requirement for successful e-Governance.
- ❖ A Quality Model based on International Standards is required.
- Quality model (ISO 25000) describes key Quality Characteristics like Functionality, Security, Reliability, Usability, Performance and Document Quality.
- Testing is done in a controlled laboratory environment and latest test tools to ensure that results are reliable, repeatable & reproducible.
- Test Reports are analysed and Software Quality Evaluation is done to find out Quality score



Conformity Assessment

Vision

To ensure transparent, cost effective, secure and citizen centric eGovernance services for the common man through evaluation by an independent Body, following global best practices

Confidence in invisible Government

❖ The results of conformity assessment are based on objective evaluation with verifiable evidences. The results are reliable, repeatable and reproducible. Claims based on conformity assessment that solution meets the requirements of "Contract" are expected to inspire greater confidence among stakeholders then claims without such backing

Viewing Reality

Conformity Assessment Framework constitutes a way of viewing reality through a ratio of Compliances and requirements

Business of Confidence- what is required?

- ❖ Sound Eco-System where buyers (Government Departments), Sellers (Solution Providers) and Conformity Assessment Bodies (Software Test Laboratories) work together in a disciplined way to achieve eGovernance objectives and timelines
- ❖ This requires Software Test Laboratories are dependable which comes from implementing a domain specific laboratory "Quality Management System"
- ❖ ISO/IEC 17025 describes requirements for this Quality Management System

Laboratory Quality Management System (Contd.) It has two components: - Management Requirements **Technical Requirements** Management Requirements ☐ Organisation Quality System ■ Document Control ☐ Review of Request, Tenders and Contracts ☐ Subcontracting of Tests and Calibrations □ Purchasing Services and Supplies ■ Service to the Clients □ Complaints ■ Control of Non-conforming Testing ■ Improvements □ Corrective Actions Preventive Actions ☐ Control of Records ■ Internal Audits ■ Management Reviews

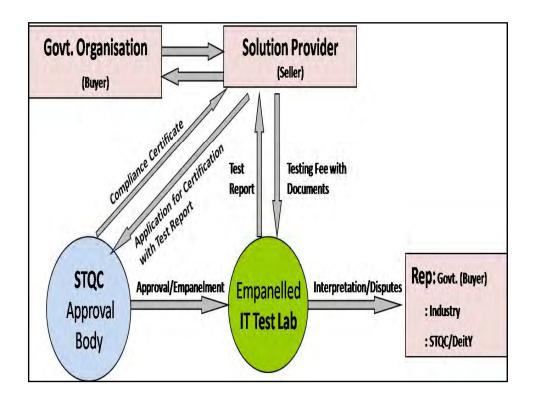
Laboratory Quality Management System Technical Requirements Personnel Accommodation and Environmental Conditions Test and Calibration Methods and Method Validation Equipment Measurement Traceability Sampling Handling of Test and Calibration Items Assuring the Quality of Test and Calibration Results Reporting the Results



Approval Scheme
For
Information Technology Testing Laboratory

WHY: CONFORMITY ASSESSMENT SCHEME

- No accredited agency for benchmarking & certifications in India
- Solution providers approach STQC labs for evaluation and certification of e Gov products
- Information technology systems and applications are more complex and domain/technology intensive.
- Managing Testing/audit of mission critical e-Governance solutions is very difficult: Collaboration, Churning, Coordination and Cycle
- STQC labs not able to meet growing demand
- Need for "Conformity Assessment Scheme"
- Look for Model for conformance testing and certification programs



Conformity Assessment

Approval Process for Software Test Houses

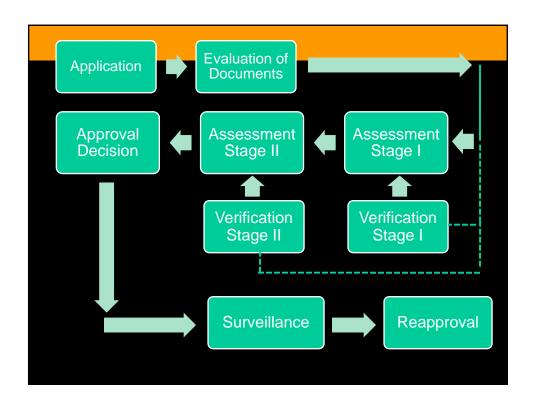
PROCESS

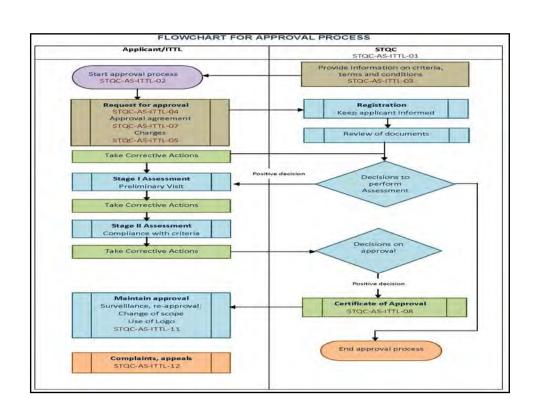
- · STQC will be the nodal agency managing the Approval Scheme
- STQC Approval body will operate as per ISO/IEC 17011: 2004 standard "Conformity Assessment-General requirement for accreditation bodies accrediting conformity assessment bodies"
- IT Testing laboratory will be assessed as per ISO/IEC 17025: 2005 General requirements for the competence of testing and calibration laboratories
- Test Management activity will be assessed as per ISO/IEC 25051 (ISO/IEC 29119 under development)
- STQC IT Bangalore as nodal agency for Technical operations of approval scheme

7/20/20,

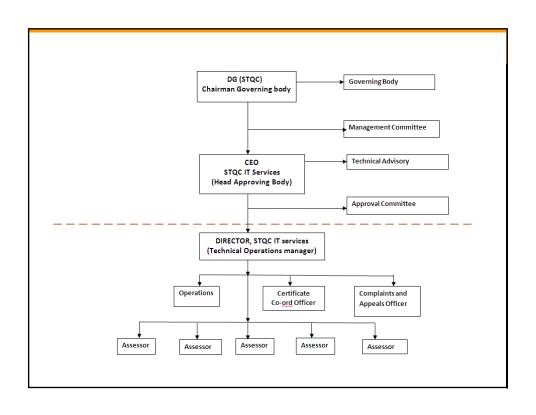
IT BANGALORE







Conformity Assessment Certification Body Organisation



SCOPE OF APPROVAL

■ Main disciplines:

- Software conformance Testing
- System conformance Testing
- Network Testing

Functional Testing: Interoperability Testing

Portability Testing **Application Security**

Testing

Performance Testing Network Testing (

Security)

Network Testing (**Usability Testing**

Performance)

Rortability Testing (Application) Code Review IT BANGALORE

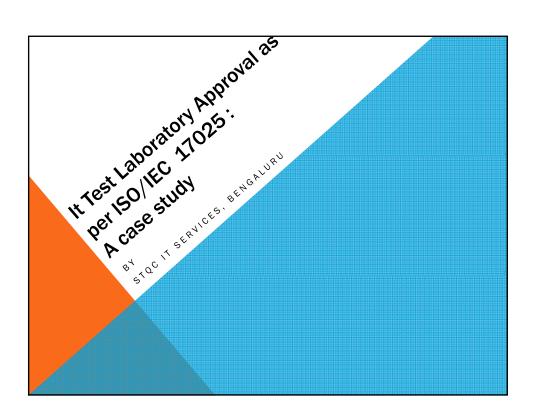
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DOCUMENT STRUCTURE				
SI. No.	Section No.	Versi on	Control Document Name	Status
1	STQC-AS- ITTL-01	1.0	Master List of Documents	complete d
2	STQC-AS- ITTL-02	1.0	Approval Scheme	by 28/2/13
3	STQC-AS- ITTL-03	1.0	Approval Criteria of IT Testing Laboratories	complete d
4	STQC-AS- ITTL-04	1.0	Application form for approval of IT Testing Laboratory	complete d
5	STQC-AS- ITTL-05	1.0	Schedule of Charges	complete d
6	STQC-AS- ITTL-06	1.0	Assessment Forms	complete d
7	STQC-AS- ITTL-07	1.0	Approval Agreement	complete d
8	STQC-AS- ITTL-08	1.0	Certificate of Approval of IT Test Laboratory	complete d
9	STQC-AS-	1.0	Check List for Approval of independent Testing	complete

Tool Kit

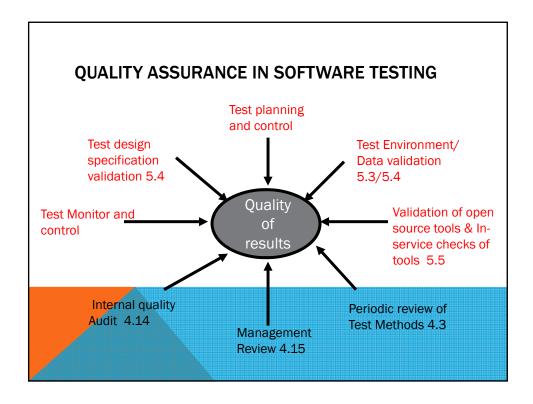
- Training Programmes
- Workshops
- Exposure to validated Procedures





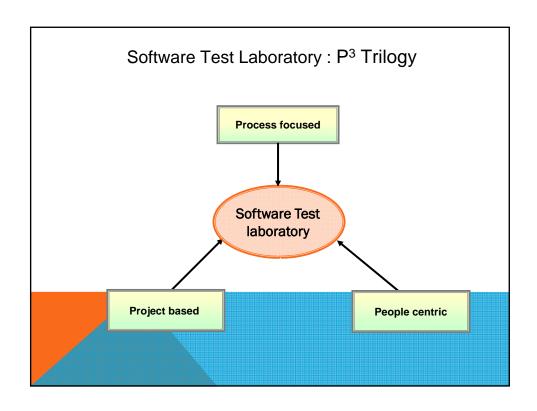
Software Test Laboratory : A2LA Accreditation: Road Map

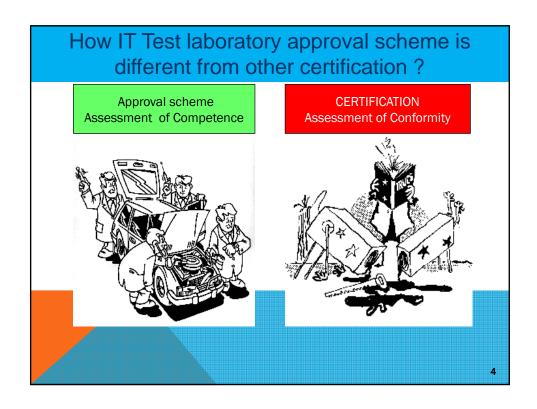
- > Scope of Operation
- > Organization structure
- > Laboratory Infrastructure Development
- > Capability and Competence Development
 - Training and Personnel certification
 - Identification of Test processes, Testing techniques, Test methods etc in line with ISO standards
 - Mapping of Test process and Laboratory Good Practices with ISO/IEC 17025
 - Establishment and Implementation of Test processes
 - Identification Test documentation requirements
 - IQA and Management Review
- Final assessment and Accreditation

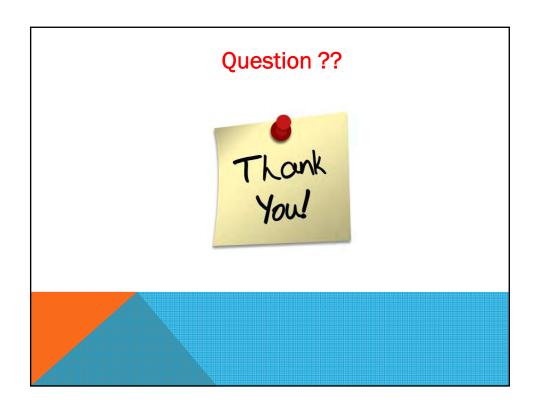


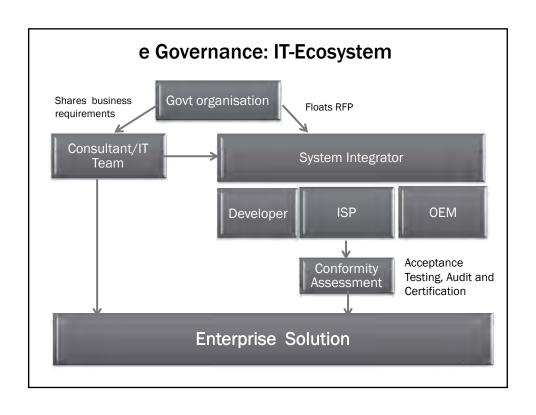
SOFTWARE TEST LABORATORY: FEW TIPS

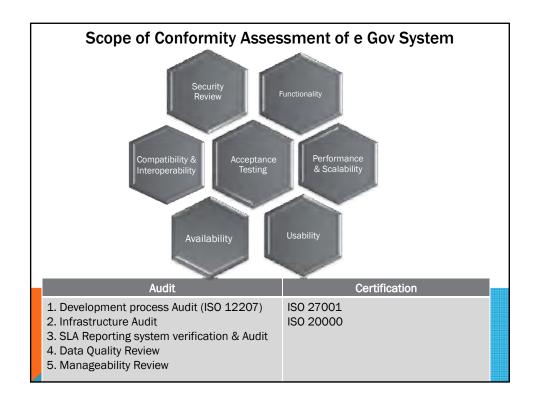
- Generally all Software applications are unique and ridden with complex business rules and algorithms.
- Each software product will entail different types of testing viz Functionality, performance, security, usability, compatibility etc.
- Most of the Technical elements of ISO/IEC 17025 are elaborated and supported with best practices in ISO/IEC Standards
- Technical challenges :
 - Domain and context understanding
 - Test coverage
 - Test environment/data creation
 - Test tool validation
- · Other Issues:
 - Schedule demands
 - Customer/ user expectations
 - Availability of Test Environment infrastructure
 - Scope of Project
 - Criticality of the project
 - Adequacy of Requirement document

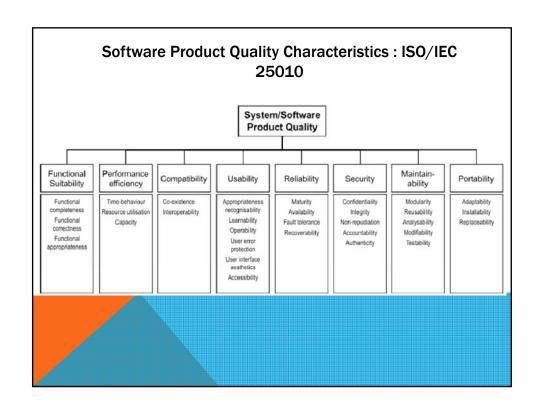




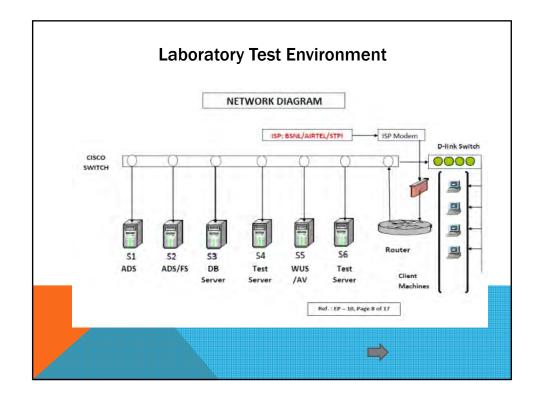








Scope of Software Test laboratory A2LA Certificate Number: 2686.01 Unit / System Tested Activity Reference Standard / Method Software Applications & Conformity Evaluation ISO/IEC 25051; EP-10 ISO/IEC 25051; EP-11 Systems* Functionality Testing Performance / Efficiency ISO/IEC 25051; EP-12 Testing Usability Testing & Heuristic ISO/IEC 9241, ISO IEC 9126-2; EP-21 Evaluation (Human - System Interaction Analysis) Static Code Review, Inspection IEEE 1028, IEEE 1044; & Maintainability Analysis ISO/IEC 9126; EP-22 Web Applications & Computer OWASP Testing Guide; EP-19 Security Testing Networks* Vulnerability Assessment, CIS; NIST SP 800-115; EP-24 Non-Destructive Penetration Testing *This laboratory meets A2LA R104 - General Requirements: Accreditation of Field Testing and Field Calibration Laboratories for these tests.



Software Testing related ISO/IEC & IEEE standards

- The following ISO/IEC & IEEE International standards are referred that
 provides a consistent set of definitions, processes, procedures, techniques &
 best practices for software testing
 - > ISO/IEC 29119 : Software and Systems engineering Software Testing

Part 1: Concepts and definitions

Part 2: Test process
Part 3: Test Documentation
Part 4: Test Techniques

(BS 7925-1:1998, Software testing - Vocabulary

BS 7925-2:1998, Software testing - Software component testing)

- ▶ IEEE Std 1028-2008, IEEE Standard for Software Reviews and Audits
- ISO/IEC 25010: Systems and software engineering —Systems and software Quality Requirements and Evaluation (SQuaRE) — System and software quality models
- ISO/IEC 25051: (Software Product Quality Requirements and Evaluation: Requirement for quality of Commercial Off-The-Shelf(COTS) software product and instruction for testing
- ISO/IEC 12207: Systems and software engineering Software life cycle processes
- FileEE Std 1012-2012, IEEE Standard for Software Verification and Validation

Software Testing related ISO/IEC & IEEE standards

- ISO/IEC 15408: Information technology Security techniques Evaluation criteria for IT security
- ISO 9241: Usability stds Ergonomics of Human system interaction. Usability Testing & Heuristic Evaluation (Human System Interaction Analysis
- ISO/IEC 23026:2006 Software Engineering Recommended Practice for the Internet Website Engineering, Web Site Management, and Web Site Life Cycle
- > NIC guidelines on Indian government websites
- OWASP (top 10 issues) for Application security Testing
- > CIS standard for Vulnerability Assessment
- NIST 800-115: Technical guide to Information Security Testing and Assessment

PROCESS MODEL Organizational Test process Test Management process

Dynamic Test process

ISO/IEC 29119: MULTI LAYER MODEL TEST PROCESS REQUIREMENTS Organizational Test process Test Policy Test Management Processes Test Planning Process Test Monitoring and Control Process Process

Dynamic Test processes

Test Execution

Process

Test Env set up

and Maintenance

Process

Test Incident

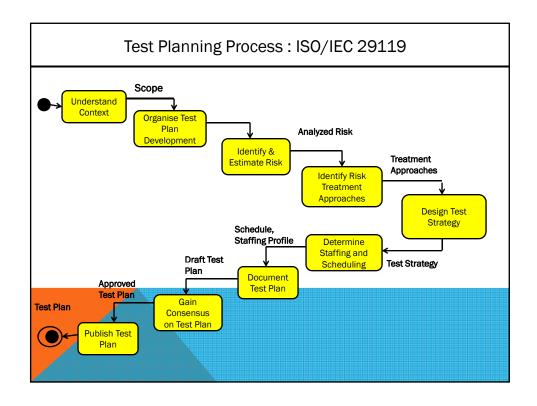
reporting

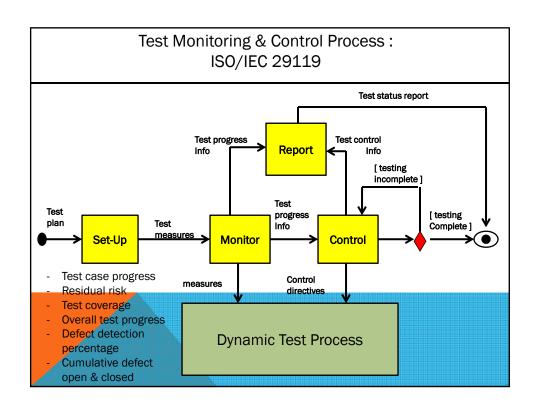
Process

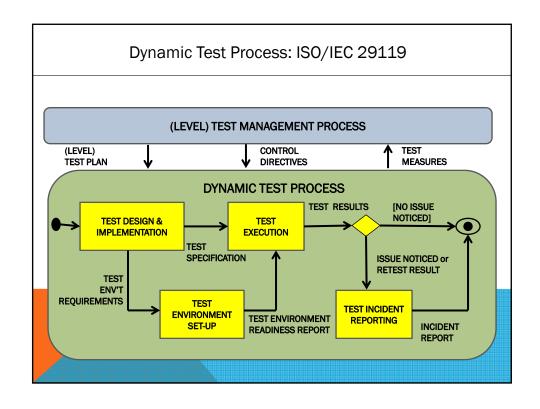
Test Design &

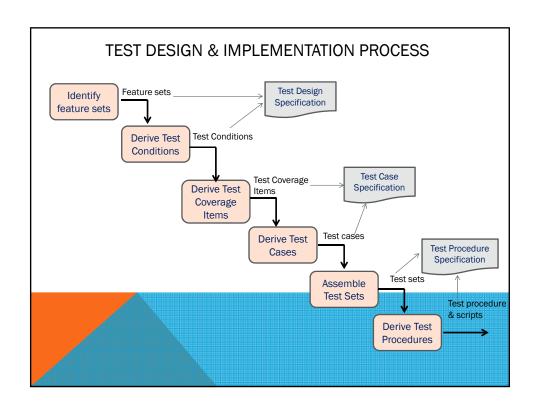
Implementation

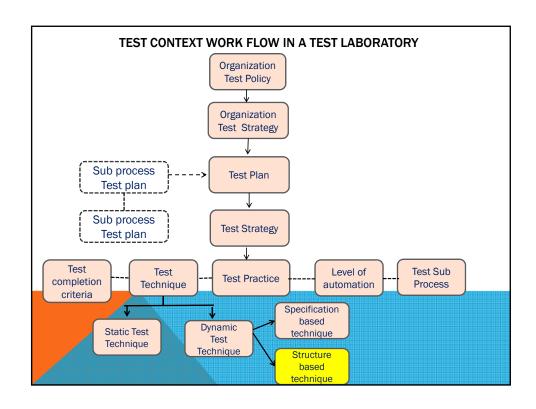
Process











Executive Procedures			
SI No.	EP NO.	Name of the procedure	ISO/IEC 17025:2005 Applicable Reference
1	EP 01	Placement of Personnel	5.2
2	EP 02	Control of documents	4.3
3	EP 03	Operation of Customer Services	4.4
4	EP 04	Internal Quality Audit	4.14
5	EP 05	Management Review	4.15
6	EP 06	Training	5.2
7	EP 07	Quality Assurance	5.9, 4.10
8	EP 08	Configuration Management	4.3, 4.13,
9	EP 09	Handling Customer Complaint and Feedback	4.7, 4.8
10	EP 10	Operation of Software Test Laboratory	5.3,5.5,5.6, 5.10
11	EP 11	Software Functional Testing	5.4
12	EP 12	Software Performance Testing	5.4
13	EP 13	Software Compatibility Testing	5.4
14	EP 14	Acceptance Testing of Software Application	5.4
15	EP 15	Software Product Evaluation	5.4
16	EP 16	Approval of Sub Contractors	4.5
17	EP 17	Control of non conforming work, and implementation of Corrective actions and preventive action	4.9,4.11,4.12
18	EP 18	Execution of on Site Testing	5.4
19	EP 19	Software Application Security Testing	5.4
20	EP 20	Procedure on Security practices in the Laboratory	5.4.7
21	EP 21	Procedure on Usability Testing	5.4
22	EP 22	Procedure for conduct of software code review	5.4

Test Methods					
SI No.	EP NO.	Title of Test Methods			
1	TM 01	Designing and Execution of test cases using Equivalence Partitioning			
2	TM 02	Designing and Execution of test cases using Boundary Value Analysis			
3	TM 03	Designing and Execution of test cases using State Transition Strategy			
4	TM 04	Designing and Execution of test cases using Cause Effect Graph			
5	TM 05	Designing and Execution of test cases using Orthogonal Array Testing Strategy			
6	TM 06	Preparation of Software Test Plan			
7	TM 07	Procedure for Anomaly Logging and Reporting			
8	TM 08	Procedure for Software Document review			
9	TM 09	Procedure for Test Report Preparation			
10	TM 10	Preparation of Software Evaluation Plan			
11	TM 11	Procedure on Regression Testing			
4					

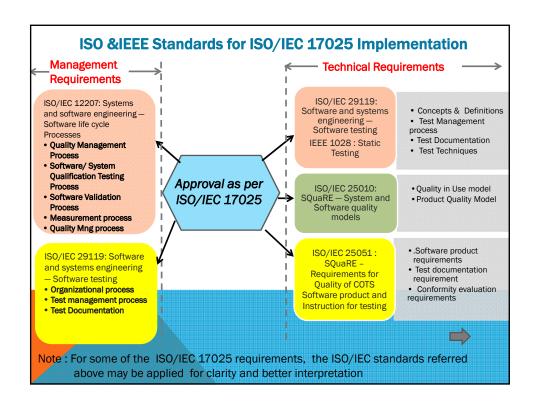
TEST DOCUMENTATION REQUIREMENT: ISO/IEC 29119

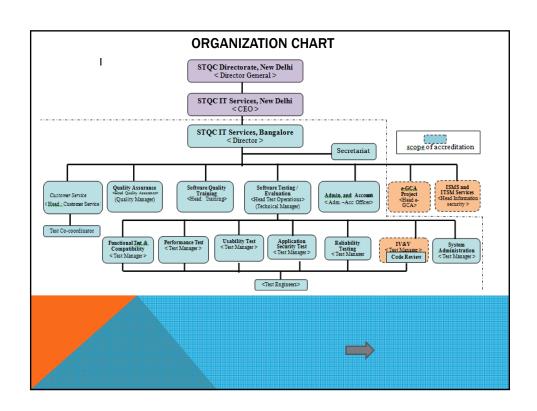
- Organizational test documentation
 - Test policy
 - Organizational Test strategy
- Test Management Processes Documentation
 Test Plan

 - Test Status Report
 - Test Completion Report
- Dynamic Test Processes Documentation
 - Test Design Specification;
 - Test Case Specification;
 - Test Procedure Specification.
 - Test Data Readiness Report
 - Test Environment Readiness Report
 - Test Execution Log.
 - Incident Report
- Test artifacts are placed under Configuration Management including Requirement
- Access rights to test artifacts is controlled
- Test artifacts such as Test Plan, Test cases, test log etc. is periodically reviewed at regular intervals and audit trial is established

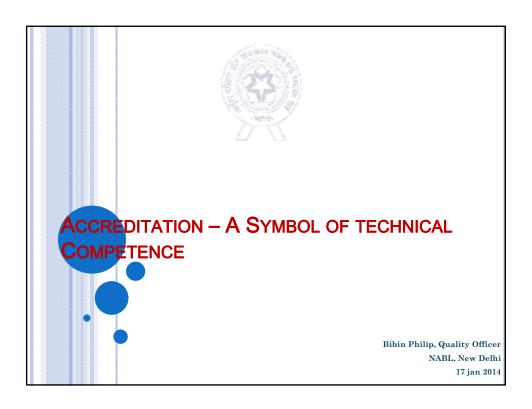
 \Rightarrow

Back up of test artifacts are taken at regular intervals





Designation	Job Profile	Academic qualification	Experience in years
est Engineer	Developing test design specifications, test cases, test suits etc.	B Tech, B E, MCA or equivalent Diploma in IT/CSc/ Electronics /Communication	- min. 2 years;
est Lead / est Manager	Developing Test Plan, Test Methods, review of test design specifications, test cases, test suits etc.	B Tech, B E, MCA or equivalent, Diploma in IT/CSc/ Electronics / Communication CSTM/ISTQB certification or equivalent is desirable;	min. 2 years min. 5 years;
echnical Manager/ est Strategist	Overall Responsibility of Technical operations Establishes, and ensures conformance to, the Organizational Test Process	B Tech, B E, MCA or equivalent, Diploma in IT/CSc/ Electronics /Communication; CSTM/ISTQB certification or equivalent is desirable,	min. 5 years
Quality Manager	Overall Responsibility of QMS activities of laboratory	B Tech, B E, MCA or equivalent, Diploma in IT/CSc/ Electronics /Communication	min. 5 years min. 10 years



OVERVIEW



- Terminologies
- Laboratory Accreditation or ISO 9001 Certification
- Why Use an Accredited Laboratory?
- Why Become an Accredited Laboratory?
- Benefits of Accreditation
- International Scenario of Laboratory Accreditation
- NABL- The National Accreditation Board

CERTIFICATION



Procedure by which a third party gives a written assurance that a product, process or service conforms to specified requirements.

(ISO/IEC 17000: 2004)

Assures the customer that the organisation has in place an effective quality or environmental management system

Does not confer technical credibility of the test result







ACCREDITATION



Third-party attestation related to a conformity assessment body conveying formal demonstration of its competence to carry out specific conformity assessment tasks

(ISO/IEC 17000: 2004)

Assures the customer that the procedures and test results are technically valid

Recognizes the technical competence of laboratory staff

•Endorses that the laboratory operates the management system effectively







LABORATORY ACCREDITATION OR ISO 9001 CERTIFICATION?



- This standard (ISO 9001) is widely used in manufacturing and service organisations to evaluate their system for managing the quality of their product or service.
- Whilst effective as a management evaluation tool, ISO 9001 does not evaluate the technical competence of a supplier. This means that the evaluation of a supplier against ISO 9001 does not assure you or your customers that the test, or calibration data are accurate and reliable.

Accreditation is more desirable for a laboratory

WHY USE AN ACCREDITED LABORATORY?

• What should you consider when choosing a laboratory? When selecting a laboratory to fulfil your testing, calibration or measurement needs, you need to be sure that they can supply you with accurate and reliable results.

The technical competence of a laboratory depends on a number of factors including:

- the qualifications, training and experience of the staff
- the right equipment properly calibrated and maintained
- adequate quality assurance procedures
- proper sampling practices
- appropriate testing procedures
- valid test methods
- traceability of measurements to national/international standards
- accurate recording and reporting procedures
- suitable testing facilities



WHY IS A LABORATORY'S TECHNICAL COMPETENCE SO CRITICAL TO YOU AS A MANUFACTURER, SUPPLIER, EXPORTER OR CUSTOMER?

- Minimise risk
- Avoid expensive retesting
- o Enhance your customers' confidence
- Reduce costs and improve acceptance of your goods overseas

Why become an Accredited Laboratory?

- A recognition of testing competence
- A benchmark for performance
- A marketing advantage?
- International recognition for your laboratory

Why become an Accredited Laboratory?

- ⇒Better control of laboratory operations
- ⇒Improves staff confidence and enhances business
- ⇒Enhanced customer confidence and satisfaction
- ⇒Reliability of data for R&D
- Insurance companies can rely on test results
- ⇒Ensures better support in the event of legal cases
- ⇒Provides traceability in measurements to national standards
- ⇒Provides global equivalence
- ⇒Saving of time & money due to re-testing

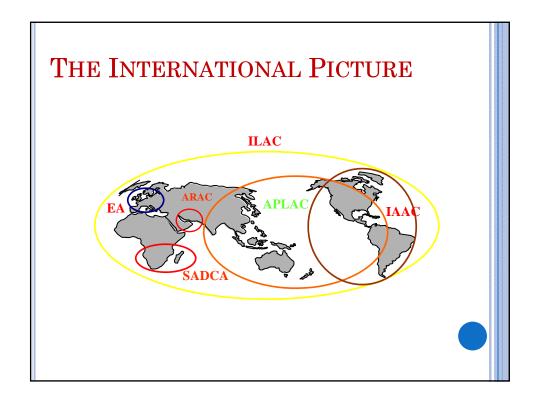


LABORATORY ACCREDITATION- MILESTONES

War time test house scheme	Australia	1943
NATA	Australia	1946
Telarc	New Zealand	1973
STP	Denmark	1973
NVLAP	USA	1976
ILAC - International Conference	1977	
A2LA	USA	1978
ISO/IEC Guide 25 - Requireme	1978	
RNE	France	1979
CSCP	Canada	1980
NCTCF	India	1981
NAMAS	UK	1981
HOKLAS	Hong Kong	1985
EA - European Co-operation	1987	
APLAC - Asia Pacific Co-opera	1992	
NABL	India	1992

MRA DEVELOPMENTS

- > Australia New Zealand Laboratory MRA -1981
- > Australia and New Zealand with UK, Sweden, Netherlands.
- > EA multilateral MRAs (Australia, NZ, HK, Singapore, USA)
- > ILAC multilateral MRA joining EA and APLAC and other regions 1996
- > APLAC multilateral MRA 1997

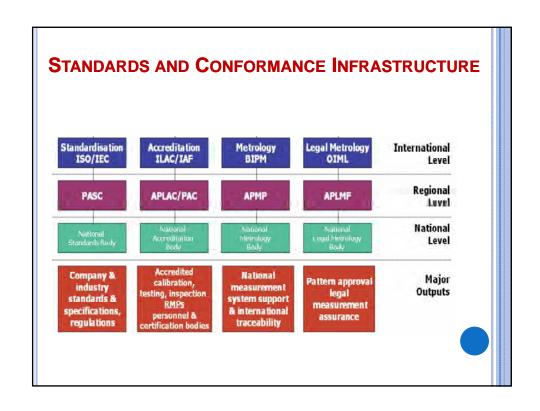


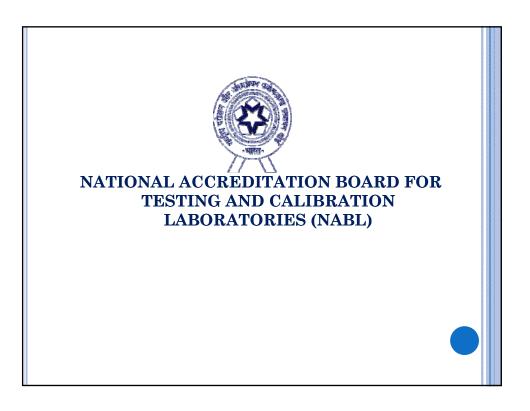
GLOBAL ACCEPTANCE OF REPORTS

MRA facilitate acceptance of test/ calibration results between economies which MRA partners represent. \

Test reports issued by an NABL accredited laboratory is considered equivalent to those issued by ILAC/APLAC MRA partners.

Global MRA has network of ~40,000 Accredited labs from 60 economies.





NABL



- ➤NABL is an autonomous body under the Department of Science & Technology (DST), Govt. of India and is registered under the Societies Act, 1860.
- NABL has been established with the objective to provide Government, Industry Associations and Industry in general with a scheme for third-party assessment of the quality and technical competence of testing and calibration laboratories.

NABL



OBJECTIVES OF NABL

- Promote & Implement Accreditation system for laboratories
- Provide timely accreditation services to laboratories
- Create awareness on Laboratory Accreditation
- Maintain data-base for Assessors & Monitor Performance
- Undertake training programs for Assessors & NABL Staff
- Operate mechanisms for Complaints and Appeals
- Maintain linkages with ILAC & APLAC through meetings
- Promote MRA with other ABs for acceptance of test report

NABL



NABL grants accreditation

- to testing & calibration laboratories as per ISO/IEC 17025
- to medical laboratories as per ISO 15189.
- To proficiency testing providers as per ISO/IEC 17043
- > NABL operates its own system as per ISO/IEC 17011.

ACCREDITATION SCOPE OF NABL

Testing Laboratories

- Biological
- Chemical
- Electrical
- Electronics
- Fluid-Flow
- Mechanical
- Non-Destructive Testing
- Optical
- Photometry
- o Radiological
- Thermal

Calibration Laboratories

- Electro-Technical
- Mechanical
- Radiological
- Thermal
- Optical
- Fluid-Flow



ACCREDITATION SCOPE OF NABLE

Medical Laboratories

- Clinical Biochemistry
- Clinical Pathology
- Haematology
- Microbiology
- Serology
- Histopathology
- Cytopathology
- Genetics
- Nuclear Medicine (In-vitro Only)

Proficiency Testing Providers

- Testing
- Calibration
- Medical
- Inspection

NABL

RECOGNITION BY GOVERNMENT & REGULATORS

- > CGHS and Government Hospital Laboratories
- > Bureau of Energy Efficiency (BEE)
- > Food Safety and Standards Authority of India
- > Central Electricity Authority (CEA)
- Agricultural & Processed Food Products Export Dev. Authority (APEDA)
- > Telecom Engineering Centre (TEC)
- > Central Pollution Control Boards (CPCB)
- > Export Inspection Agency (EIA)
- > Tea Board, Coffee Board, Spices Board
- > Judiciary also relies on accredited forensic laboratori

INTERNATIONAL RECOGNITION OF NABL

- NABL is signatory to Asia Pacific Laboratory Accreditation Co-operation (APLAC) Mutual Recognition Arrangements (MRA) and International Laboratory Accreditation Co-operation (ILAC) MRA since 2000.
- → NABL was evaluated by APLAC in 2000,2004,2008 and 2012 and current MRA valid till 2016







INTERNATIONAL RECOGNITION OF NABL

- NABL has been recognized as approved accreditation body by Environmental Protection Agency (EPA) USA for it Energy Star labeling scheme
- NABL has also been recognized by **The LED Lighting Facts**, **USA** as approved accreditation body. NABL is the first Accreditation Body outside USA to get such a recognition.







NABL ACCREDITED LABORATORIES

Govt. Department Laboratories (approx 200)

- 1. Department of Atomic Energy
- 2. Ministry of Agriculture
- 3. Ministry of Chemicals & Fertilizers
- 4. Ministry of Commerce & Industry
- 5. Ministry of Communications & Information Technology
- 6. Ministry of Consumer Affairs & Public Distribution
- 7. Ministry of Defence
- 8. Ministry of Environment & Forests
- Ministry of Heavy Industries & Public Enterprises
- 10. Ministry of Home Affairs
- 11. Ministry of Petroleum & Natural Gas
- 12. Ministry of Power
- 13. Ministry of Small Scale Industries
- 14. Ministry of Textiles
- Council of Scientific and Industrial Research (CSIR)

CUSTOMERS

Private Organization Laboratories

- 1. SGS India Pvt. Ltd.,
- 2. TUV Rehinland Pvt Ltd
- 3. BVCPS India Pvt. Ltd
- 4. Intertek India Pvt. Ltd,
- 5. UL India Pvt. Ltd.
- 6. UltraTech Cement Limited.
- 7. Birla Cement Works
- 8. Larsen and Toubro Limited
- 9. Raymond Limited
- 10. Godfrey Philips India Limited
- 11. Sundaram Auto Components Ltd
- 12. Whirlpool of India Limited
- 13. Blue Star Limited, Chennai
- 14. Mahindra & Mahindra Ltd

CUSTOMERS

Private Organization Laboratories

- Torrent Power Ltd.
- Maruti Udyog Ltd.
- TATA Power/TISCO/TATA Refractories / TATA Steel
- Reliance Industries Ltd.
- HCL Itd.
- GE India Technology Centre
- TUV Sud South Asia Pvt. Ltd.
- Hindustan Coca-Cola Beverages Pvt. Ltd.
- ITC R&D Centre
- Robert Bosch Engineering Solutions Ltd
- MICO Application Centre (MICO Bosch Group)
- MRF R&D Corporate Technical

CUSTOMERS

Private Organization Laboratories

- SRL Ranbaxy Limited
- Dr. Lal Path Labs Pvt., Ltd.
- Max Labs 24X7, Max Super Speciality Hospital
- Apollo Gleneagles Hospitals
- Wockhardt Laboratory Services
- TATA Memorial Hospital
- B. M. Birla Heart Research Centre
- Reliance Life Sciences Pvt. Ltd.
- Manipal Acunova Central Reference Laboratory
- Sitaram Bhartia Institute of Science & Research
- Sir Ganga Ram Hospital, Clinical Laboratory Services
- Metropolis Health Services (India) private Limited

CUSTOMERS

Cross Border Laboratory Accreditation

- 1. SGS Bangladesh Ltd, Dhaka, Bangladesh
- 2. Bureau Veritas Consumer Products Services Ltd., Dhaka, Bangladesh
- 3. Textile Training & Services Centre Textile Testing Laboratory, Sri Lanka
- 4. Bureau Veritas Consumer Products Services (Lanka) Ltd., Sri Lanka
- 5. SGS Lanka Pvt. Ltd, Colombo, Sri Lanka
- 6. Zest Laboratories Pvt. Ltd. Nepal
- 7. TUV SUD Bangladesh Pvt. Ltd., Bangladesh
- 8. CFL, Department of Food Technology & Quality Control, Nepal
- 9. TUV Rheinland Bangladesh Pvt. Ltd. Bangladesh



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