

INF5180: Software Product- and Process Improvement in Systems Development

Part 09: Process Assessment



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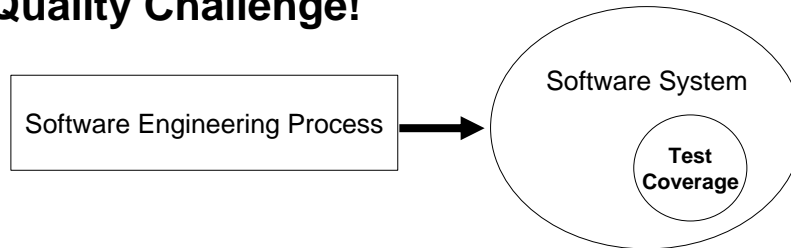
Spring 2010

Contents

- CMM(I) History and Overview
- Structure
 - Specific and generic goals
 - Specific and generic practices
- Process Areas
- Evaluation
- Continuous Process Improvement
- Dissemination and Results



The Quality Challenge!



"The quality of a system is highly influenced by the quality of the process used to acquire, develop, and maintain it." (W.Humphrey)



The Software Business ...

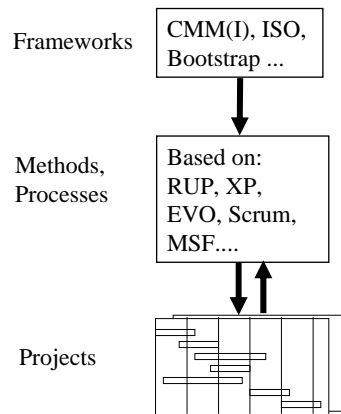
- Problems with estimation
 - Budget overruns
 - Time overruns
- Problems with quality
 - Struggle with defect correction instead of avoidance
 - Struggle with unsatisfied customers
- No well-defined process
 - Starting "from scratch" in every project
 - Good practices are sacrificed under stress
- Critical aspects
 - Relying on "heroes"
 - Overtime
 - Fire-fighting



... and this is even more the case for system development



How to Sustain Good Practice?



Maturity/quality frameworks support

- evaluating improvement needs and achievements
- organizational change
- cultural change
- knowledge creation
- measurement

Much good practice is prescribed in explicit process models. But you should **choose right** and **adapt** to your needs

It is important to identify and learn about good practices from within the projects. Update the process model after each project!



Origins of CMM*

- DoD** decided in the 80s to do something about everything that went wrong in its expensive software projects (often involving suppliers).
 - Ada didn't solve the problems (as many had thought/hoped)
 - Appraisals showed that there was a management problem
- Watts Humphrey left IBM for SEI and began developing CMM in 1986
- *Managing the Software Process* by W. Humphrey published in 1989
- Version 1.1 published in 1993 – is still the most used model
- CMMI first published in 1999, version 1.2 published in 2006.

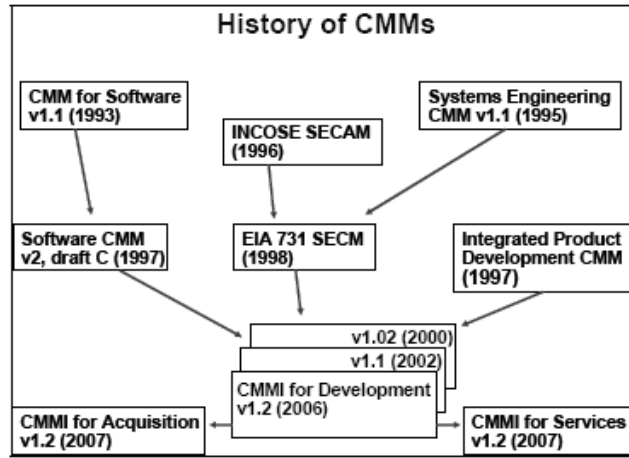
CMM = Capability Maturity Model
DoD** = Department of Defense



History

Source: CMMI® for Development, Version 1.2 (CMMI-DEV, V1.2), CMU/SEI-2006-TR-008, ESC-TR-2006-008, August 2006

<http://www.sei.cmu.edu/cmmi/>



Page 7

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CMMI Family

- 4 different models (with very small differences)
 - CMMI-SE/SW/IPPD/SS
 - CMMI-SE/SW/IPPD
 - CMMI-SE/SW
 - CMMI-SW
 - All models have a continuous and staged representation.
- Definitions:
 - SS: Supplier Sourcing
 - IPPD: Integrated Product and Process Development
 - SE: Systems Engineering
 - SW: Software Engineering

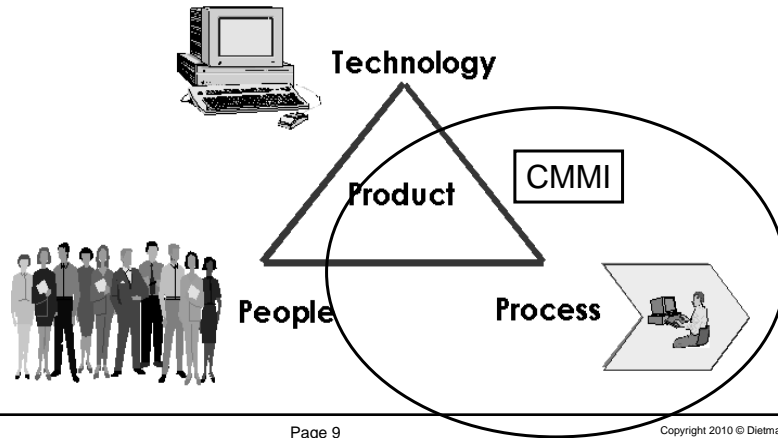
Page 8

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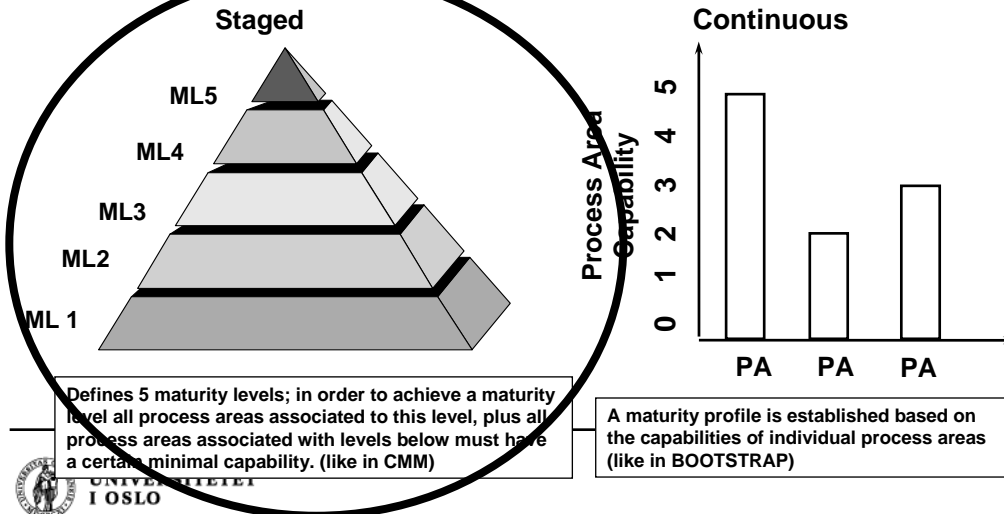


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CMMI Scope



Staged versus Continuous Representation



Levels and Process Areas (staged)

* Integrated Product/Process Development (IPPD) – add-on to the Engineering processes
 ** Acquisition – add-on to the Engineering processes



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Level	Process Areas
5 Optimizing	Causal Analysis and Resolution Organizational Innovation and Deployment
4 Quantitatively Managed	Quantitative Project Management Organizational Process Performance
3 Defined	Requirements Development Technical Solution Product Integration Verification Validation Organizational Process Focus Organizational Process Definition Organizational Training Risk Management Integrated Project Management (for IPPD*) Integrated Teaming* Integrated Supplier Management** Decision Analysis and Resolution Organizational Environment for Integration*
2 Managed	Requirements Management Project Planning Project Monitoring and Control Supplier Agreement Management Measurement and Analysis Process and Product Quality Assurance Configuration Management
1 Performed	

Page 11

CMMI Level 2 – *Managed* means ...

- Adhering to organizational policies
- Following established plans and process descriptions
- Providing adequate resources (including funding, people, and tools)
- Assigning responsibility and authority for performing the process
- Training the people performing and supporting the process
- Placing designated work products under appropriate levels of configuration management
- Identifying and involving relevant stakeholders
- Monitoring and controlling the performance of the process against the plans for performing the process and taking corrective actions
- Objectively evaluating the process, its work products, and its services for adherence to the process descriptions, standards, and procedures, and addressing noncompliance
- Reviewing the activities, status, and results of the process with higher level management, and taking corrective action

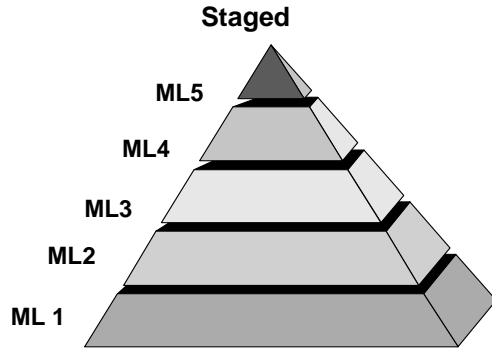
Page 12

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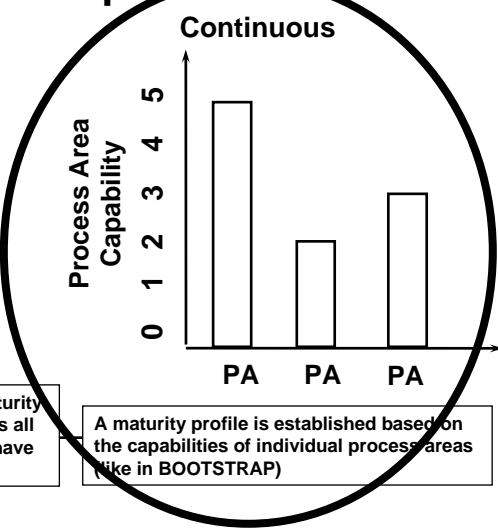


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Staged versus Continuous Representation



Defines 5 maturity levels; in order to achieve a maturity level all process areas associated to this level, plus all process areas associated with levels below must have a certain minimal capability. (like in CMM)



A maturity profile is established based on the capabilities of individual process areas (like in BOOTSTRAP)

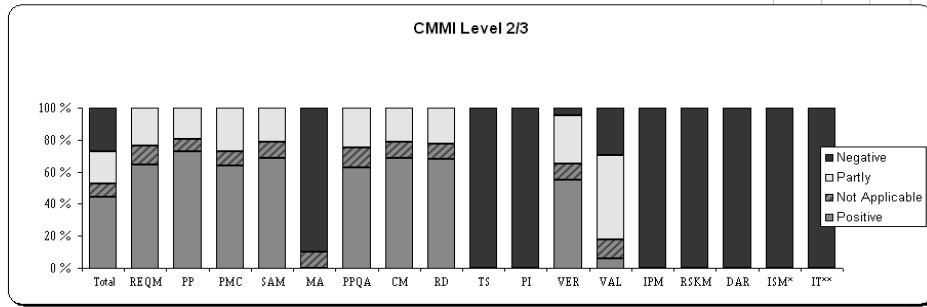


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Assessment Results (continuous)

Total CMMI Compliance: 63 %
 CMMI Level 2 compliance: 77 %

Tailoring: Use of Not Applicable: 8 %
 CMMI Level 3 compliance: 22 %



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Research Results at Simula

Headline:
That's why it goes so wrong

On average:
35% cost overrun
25% time overrun

50% of all projects are really bad

Reasons:
Underestimation of complexity
and thus
of the risks associated with
sw/system development



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COMPUTERWORLD
www.computerworld.no 20 år

Nr. 44 - Uke 42
Fredag
17. oktober
2003
21. årgang
Lesesalg kr. 40

Derfor går det så galt

Endelig er det vitenskapelig bevist: It-prosjekter blir i snitt 35 prosent dyrere og tar 25 prosent lenger tid enn planlagt. Prosjektene er nesten alltid langt mer komplekse enn antatt, hevder forskere ved Simula-senteret. – Halvparten av alle prosjekter går riktig galt. Som regel på grunn av en sterk undervurdering av risiko, sier professor Magne Jørgensen.

FOTO: ØYVIND ALSTAD

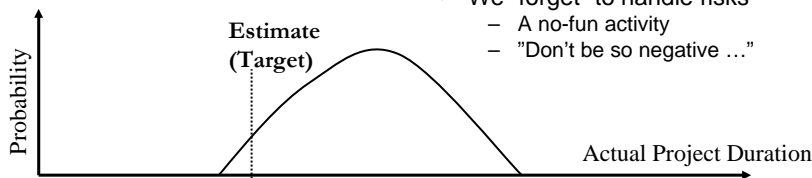
VANN OVER HODET: It-prosjekter blir alltid mer omfattende enn man tror i utgangspunktet, fastslår Kjetil Molækken og Magne Jørgensen ved Simula-senteret.

SIDE 10 OG 11

It is human to make mistakes ...

Source:
Magne Jørgensen et.al. (2003)
"Better sure than safe?
Overconfidence in judgment based software
development effort prediction intervals"

- Overestimating your own skills
 - "Everyone" is an above average driver...*
 - Study at Simula shows similar tendency for developers
- We estimate "internally"
 - Fail to look at the project from outside
 - Which experience was made with similar projects?
 - Have there been done at all similar projects before?
- We "forget" to handle risks
 - A no-fun activity
 - "Don't be so negative ..."

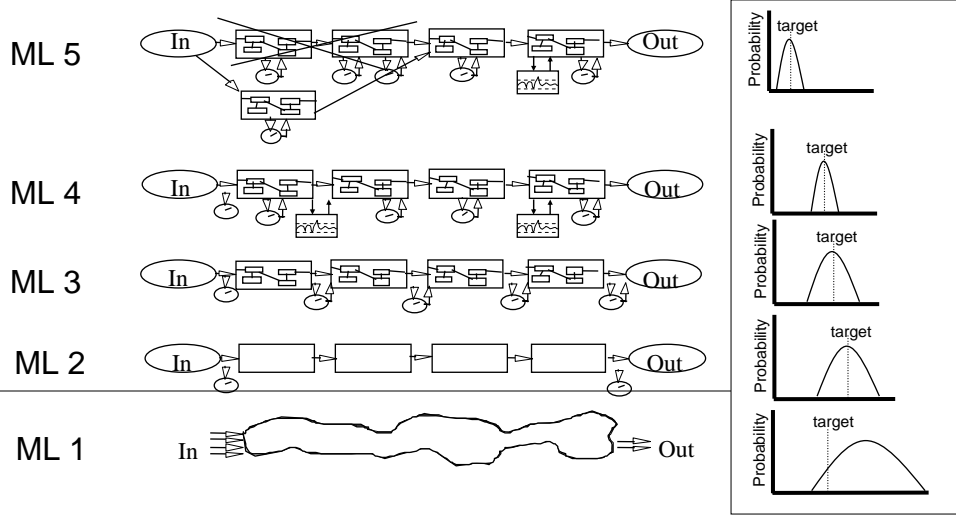


*) Applies mostly to men

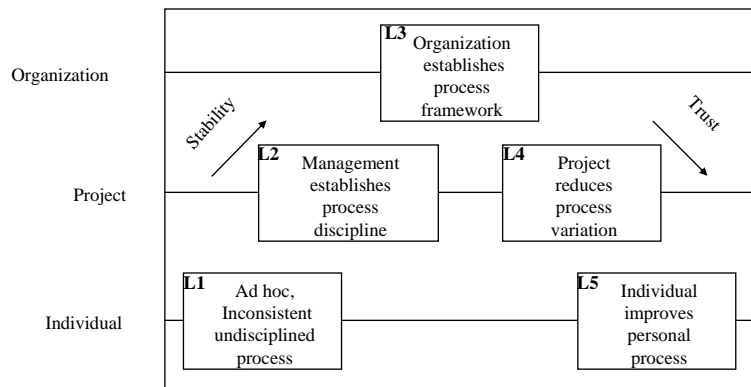


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CMMI – How much we now about our projects?



Organizational Changes

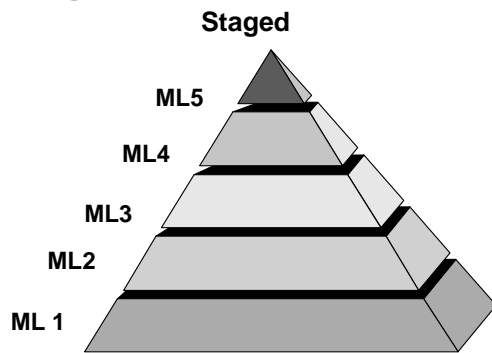




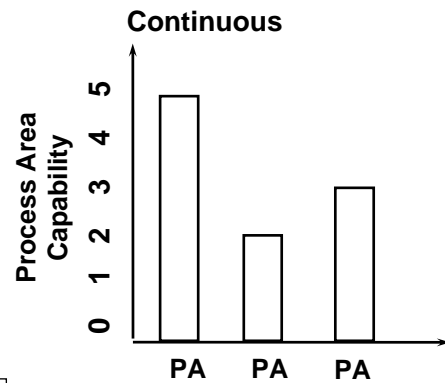
STRUCTURE



Staged or Continuous Representation?



Defines 5 maturity levels; in order to achieve a maturity level all process areas associated to this level, plus all process areas associated with levels below must have a certain minimal capability. (like in CMM)



A maturity profile is established based on the capabilities of individual process areas (like in BOOTSTRAP)

Levels and Process Areas (staged)

* Integrated Product/Process Development (IPPD) – add-on to the Engineering processes
 ** Acquisition – add-on to the Engineering processes

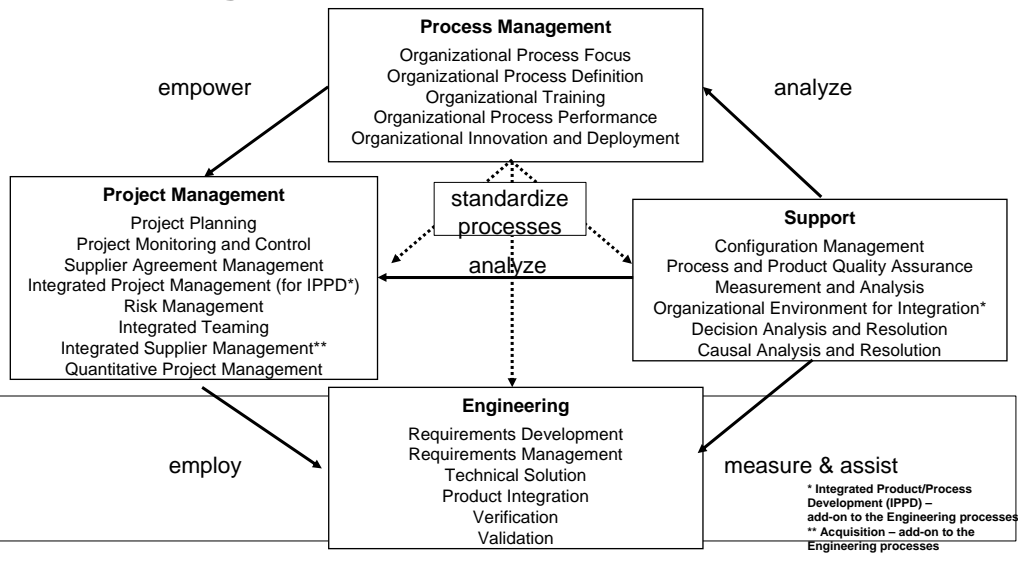


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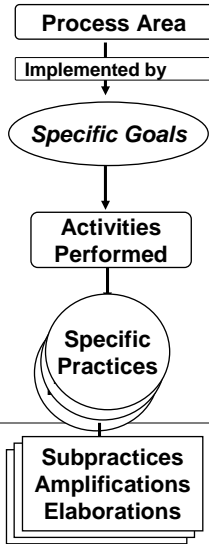
Page 21

Structuring the Process Areas (continuous)



Example: Requirement Management

The purpose of Requirements Management is to manage the requirements of the project's products and product components and to identify inconsistencies between those requirements and the project's plans and work products.



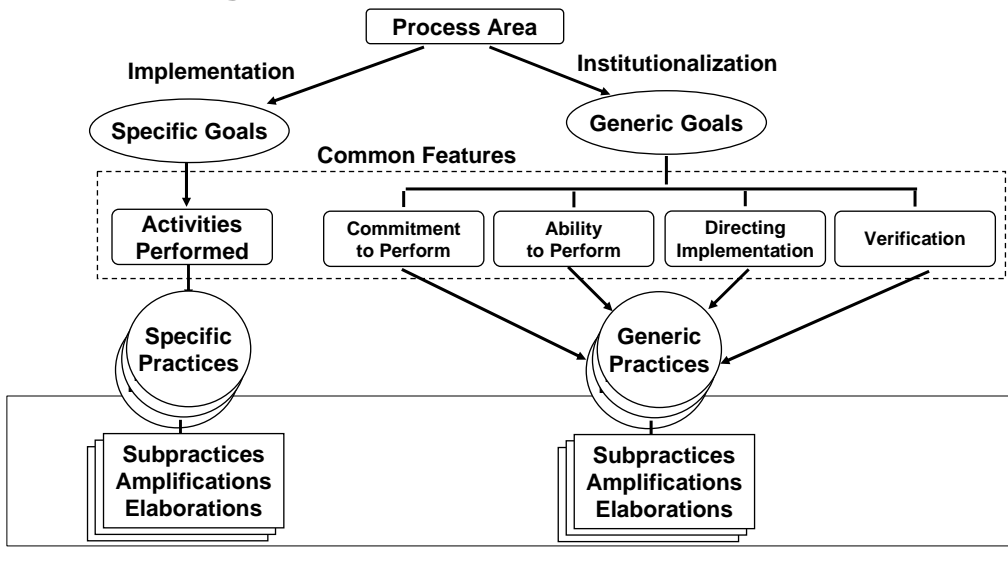
SG1: *Requirements are managed and inconsistencies with project plans and work products are identified.*

SP 1.1 Obtain an Understanding of Requirements

....

1. Establish criteria for distinguishing appropriate requirements providers.
2. Establish objective criteria for the acceptance of requirements.

Structuring the Process Areas



Specific versus Generic

- Addresses one process area
- Describes activities used to **implement** the process area
- Addresses all process areas
- Describes activities that **institutionalize** the process area

Example: Requirements Mgmt.

SG 1 Manage Requirements

- SP 1.1 Obtain an Understanding of Requirements
- SP 1.2 Obtain Commitment to Requirements
- SP 1.3 Manage Requirements Changes
- SP 1.4 Maintain Bidirectional Traceability of Requirements
- SP 1.5 Identify Inconsistencies between Project Work and Requirements

GG 1 Achieve Specific Goals

- GP 1.1 Perform Base Practices

GG 2 Institutionalize a Managed Process

- GP 2.1 Establish an Organizational Policy
- GP 2.2 Plan the Process
- GP 2.3 Provide Resources
- GP 2.4 Assign Responsibility
- GP 2.5 Train People
- GP 2.6 Manage Configurations
- GP 2.7 Identify and Involve Relevant Stakeholders
- GP 2.8 Monitor and Control the Process
- GP 2.9 Objectively Evaluate Adherence
- GP 2.10 Review Status with Higher Level Management

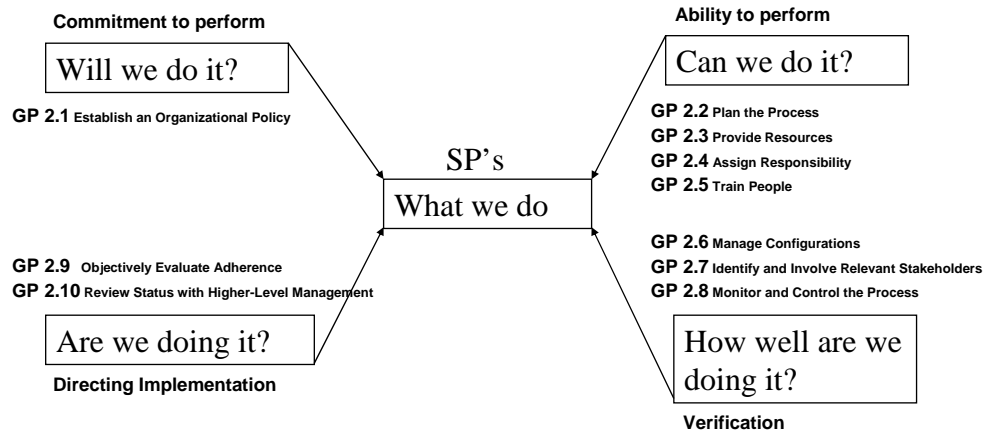
GG 3 Institutionalize a Defined Process

- GP 3.1 Establish a Defined Process
- GP 3.2 Collect Improvement Information

...



Basis for Institutionalization



Example: GP 2.2 – Plan the Process

<p>Establish and maintain the plan for performing the process.</p> <p>The purpose of this generic practice is to determine what is needed to perform the process and achieve the established objectives, to prepare a plan for performing the process, to prepare a process description, and to get agreement on the plan from relevant stakeholders. [GP104]</p> <p>Requirements for the process's specified work products and for performing the work may be derived from other requirements. In the case of a project's processes, they may come from that project's requirements management process, in the case of an organization's process, they may come from organizational sources. [GP104.N101]</p> <p>The objectives for the process may be derived from other plans (e.g., the project plans), included are objectives for the specific situation, including quality, cost, and schedule objectives. For example, an objective might be to reduce the cost of performing a process for this implementation.</p> <p>Stakeholder</p> <p>Although a generic practice imposes the practical implications on the process area, as they relate to this generic practice area, the process area may include project members, suppliers, customers, end users, and others. [FM114.HDA102.HDB104.T101]</p> <p>planning described by this generic practice as applied to the Project Planning process area typically would not be addressed by the processes associated with other process areas in the model. Therefore, the generic practice sets an expectation that the project planning process itself be planned. It is important to be aware of the extent to which this generic practice may either reinforce expectations set elsewhere in the model, or set new expectations that should be addressed. [GP104.N102]</p> <p>Establishing a plan includes documenting the plan and providing a process description. Maintaining the plan includes changing it as necessary, in response to either corrective actions or to changes in requirements and objectives for the process. [GP104.N103]</p>	<p>The Subpractices</p> <ol style="list-style-type: none"> 1. Obtain management sponsorship for performing the process. [GP104.SubP101] 2. Define and document the process description. [GP104.SubP102] 3. Define and document the plan for performing the process. 4. Review the plan with relevant stakeholders and get their agreement. [GP104.SubP104] 5. Revise the plan as necessary. [GP104.SubP105] <p>The process description, which includes relevant standards and procedures, may be included as part of the plan for performing the process or may be included in the plan by reference. [GP104.SubP102.N101]</p> <p>document, embedded in a more comprehensive multiple documents. In the case of the plan being documents, ensure that a coherent picture is preserved may be hardcopy or softcopy. [GP104.SubP103.N102]</p> <p>This includes reviewing that the planned process satisfies the applicable policies, plans, requirements, and standards to provide assurance to relevant stakeholders. [GP104.SubP104.N101]</p>
--	---

All Generic Goals

- | |
|---|
| <p>GG 1 Achieve Specific Goals [CL102.GL101]</p> <p><i>The process supports and enables achievement of the specific goals of the process area by transforming identifiable input work products to produce identifiable output work products.</i></p> |
| <p>GG 2 Institutionalize a Managed Process [CL103.GL101]</p> <p><i>The process is institutionalized as a managed process.</i></p> |
| <p>GG 3 Institutionalize a Defined Process [CL104.GL101]</p> <p><i>The process is institutionalized as a defined process.</i></p> |
| <p>GG 4 Institutionalize a Quantitatively Managed Process [CL105.GL101]</p> <p><i>The process is institutionalized as a quantitatively managed process.</i></p> |
| <p>GG 5 Institutionalize an Optimizing Process [CL106.GL101]</p> <p><i>The process is institutionalized as an optimizing process.</i></p> |



PROCESS AREAS



CMMI Level 2 – *Managed* means ...

- Adhering to organizational policies
- Following established plans and process descriptions
- Providing adequate resources (including funding, people, and tools)
- Assigning responsibility and authority for performing the process
- Training the people performing and supporting the process
- Placing designated work products under appropriate levels of configuration management
- Identifying and involving relevant stakeholders
- Monitoring and controlling the performance of the process against the plans for performing the process and taking corrective actions
- Objectively evaluating the process, its work products, and its services for adherence to the process descriptions, standards, and procedures, and addressing noncompliance
- Reviewing the activities, status, and results of the process with higher level management, and taking corrective action



CMMI Level 2

Process Areas

- Requirements Management
- Project Planning
- Project Monitoring and Control
- Supplier Agreement Management
- Measurement and Analysis
- Process and Product Quality Assurance
- Configuration Management



Requirements Management – REQM

Purpose

The purpose of Requirements Management is to manage the requirements of the project's products and product components and to identify inconsistencies between those requirements and the project's plans and work products.

SG 1 Manage Requirements

Requirements are managed and inconsistencies with project plans and work products are identified.

GG 2 Institutionalize a Managed Process

The process is institutionalized as a managed process.



Requirements Management – REQM

Practices by Goal:

GG 1 Achieve Specific Goals

SG 1 Manage Requirements

Requirements are managed and inconsistencies with project plans and work products are identified. [PA148,IG101]

SP 1.1-1 Obtain an Understanding of Requirements
Develop an understanding with the requirements providers on the meaning of the requirements. [PA148,IG101,SP101]

SP 1.2-2 Obtain Commitment to Requirements
Obtain commitment to the requirements from the project participants. [PA148,IG101,SP102]

SP 1.3-1 Manage Requirements Changes
Manage changes to the requirements as they evolve during the project. [PA148,IG101,SP103]

SP 1.4-2 Maintain Bidirectional Traceability of Requirements
Maintain bidirectional traceability among the requirements and the project plans and work products. [PA148,IG101,SP104]

SP 1.5-1 Identify Inconsistencies between Project Work and Requirements
Identify inconsistencies between the project plans and work products and the requirements. [PA148,IG101,SP105]

The process supports and enables achievement of the specific goals of the process area by transforming identifiable input work products to produce identifiable output work products.

GP 2.5 Train People

Train the people performing or supporting the process as needed. [GP107]

GP 2.6 Manage Configurations

Place designated work products of the process under appropriate levels of configuration management. [GP108]

GP 2.7 Identify and Involve Relevant Stakeholders

Identify and involve the relevant stakeholders as planned. [GP109]

GP 2.8 Monitor and Control the Process

Monitor and control the process against the plan for performing the process and take appropriate corrective action. [GP110]

GP 2.9 Objectively Evaluate Adherence

Objectively evaluate adherence of the process against its process description, standards, and procedures, and address noncompliance. [GP111]

GP 2.10 Review Status with Higher Level Management

Review the activities, status, and results of the process with higher level management and resolve issues. [GP112]

Project Planning – PP

Purpose

The purpose of Project Planning is to establish and maintain plans that define project activities.

SG 1 Establish Estimates

Estimates of project planning parameters are established and maintained.

SG 2 Develop a Project Plan

A project plan is established and maintained as the basis for managing the project.

SG 3 Obtain Commitment to the Plan

Commitments to the project plan are established and maintained.

GG1 & 2



Project Monitoring and Control – PMC

Purpose

The purpose of Project Monitoring and Control is to provide understanding into the project's progress so that appropriate corrective actions can be taken when the project's performance deviates significantly from the plan.

SG 1 Monitor Project Against Plan
Actual performance and progress of the project is monitored against the project plan.

SG 2 Manage Corrective Action to Closure
Corrective actions are managed to closure when the project's performance or results deviate significantly from the plan.

GG1 & 2



Supplier Agreement Management – SAM

Purpose

The purpose of Supplier Agreement Management is to manage the acquisition of products and services from suppliers external to the project for which there exists a formal agreement.

SG 1 Establish Supplier Agreements
Agreements with the suppliers are established and maintained.

SG 2 Satisfy Supplier Agreements
Agreements with the suppliers are satisfied by both the project and the supplier.

GG1 & 2



Measurement and Analysis – MA

Purpose

The purpose of Measurement and Analysis is to develop and sustain a measurement capability that is used to support management information needs.

SG 1 Align Measurement and Analysis Activities
Measurement objectives and practices are aligned with identified information needs and objectives.

SG 2 Provide Measurement Results
Measurement results that address identified information needs and objectives are provided.

GG1 & 2

Process and Product Quality Assurance – PPQA

Purpose

The purpose of Process and Product Quality Assurance is to provide staff and management with objective insight into the processes and associated work products.

SG 1 Objectively Evaluate Processes and Work Products
Adherence of the performed process and associated work products and services to applicable process descriptions, standards and procedures is objectively evaluated.

SG 2 Provide Objective Insight
Noncompliance issues are objectively tracked and communicated, and resolution is ensured.

GG1 & 2

Configuration Management – CM

Purpose

The purpose of Configuration Management is to establish and maintain the integrity of work products using configuration identification, configuration control, configuration status accounting, and configuration audits.

SG 1 Establish Baselines

Baselines of identified work products are established and maintained.

SG 2 Track and Control Changes

Changes to the work products under configuration management are tracked and controlled.

SG 3 Establish Integrity

Integrity of baselines is established and maintained.

GG1 & 2



CMMI Level 3 – *Defined* means ...

- Establish a **standard development process** which is
 - **Well-defined** at the organizational level
 - **in use** on a broad scale
 - the basis for all learning and storing of experience (*best practices*)
 - the starting point for special adjustments (*tailoring*)
- The organization stresses the use of the process:
 - creates process groups ("SEPG")
 - provides experience mechanism ("de-briefing", project evaluation etc)
 - links experience data to the process
 - offers training about the process
 - and ties technical training into the process
 - clearly defines interfaces between groups



CMMI Level 3

Process Areas

- Requirements Development
- Technical Solution
- Product Integration
- Verification
- Validation
- Organizational Process Focus
- Organizational Process Definition
- Organizational Training
- Integrated Project Management
- Risk Management
- Decision Analysis and Resolution



Requirements Development – RD

Purpose

The purpose of Requirements Development is to produce and analyze customer, product, and product component requirements.

SG 1 Develop Customer Requirements

Stakeholder needs, expectations, constraints, and interfaces are collected and translated into customer requirements.

SG 2 Develop Product Requirements

Customer requirements are refined and elaborated to develop product and product component requirements for the product life cycle.

SG 3 Analyze and Validate Requirements

The requirements are analyzed and validated, and a definition of required functionality is developed.

GG1-3


Technical Solution – TS

Purpose

The purpose of Technical Solution is to develop, design, and implement solutions to requirements. Solutions, designs and implementations encompass products, product components, and product related processes either singly or in combinations as appropriate.

SG 1 Select Product Component Solutions
Product or product component solutions, including applicable product related processes, are selected from alternative solutions.

SG 2 Develop the Design
Product or product component designs are developed.

SG 3 Implement the Product Design
Product components, and associated support documentation, are implemented from their designs.

GG1-3


Product Integration – PI

Purpose

The purpose of Product Integration is to assemble the product from the product components, ensure that the product, as integrated, functions properly, and deliver the product.

SG 1 Prepare for Product Integration
The strategy for conducting product integration is established and maintained.

SG 2 Ensure Interface Compatibility
The product component interfaces, both internal and external, are compatible.

SG 3 Assemble Product Components and Deliver the Product
Verified product components are assembled and the integrated, verified, and validated product is delivered.

GG1-3


Verification – VER

Purpose

The purpose of Verification is to assure that selected work products meet their specified requirements.

- SG 1 Prepare for Verification**
Preparation for verification is conducted.

- SG 2 Perform Peer Reviews**
Peer reviews are performed on selected work products.

- SG 3 Verify Selected Work Products**
Selected work products are verified against their specified requirements.

GG1-3

Validation – VAL

Purpose

The purpose of Validation is to demonstrate that a product or product component fulfills its intended use when placed in its intended environment.

- SG 1 Prepare for Validation**
Preparation for validation is conducted.

- SG 2 Validate Product or Product Components**
The product or product components are validated to ensure that they are suitable for use in their intended operating environment.

GG1-3

Organizational Process Focus – OPF

Purpose

The purpose of Organizational Process Focus is to establish and maintain an understanding of the organization's processes and process assets, and to identify, plan, and implement the organization's process improvement activities.

SG 1 Determine Process Improvement Opportunities
Strengths, weaknesses, and improvement opportunities for the organization's processes are identified periodically and as needed.

SG 2 Plan and Implement Process Improvement Activities
Improvements are planned and implemented, process assets are deployed, and process-related experiences are incorporated into the organization's process assets.

GG1-3

Organizational Process Definition – OPD

Purpose

The purpose of Organizational Process Definition is to establish and maintain a usable set of organizational process assets.

SG 1 Create Organizational Process Assets
A set of organizational process assets is available.

SG 2 Make Supporting Process Assets Available
Process assets that support the use of the organization's set of standard processes are available.

GG1-3

Organizational Training – OT

Purpose

The purpose of Organizational Training is to develop the skills and knowledge of people so they can perform their roles effectively and efficiently.

SG 1 Identify Training Needs and Make Training Available
Training to support the organization's management and technical roles is identified and made available.

SG 2 Provide Necessary Training
Training necessary for individuals to perform their roles effectively is provided.

GG1-3

Integrated Project Management – IPM

Purpose

The purpose of Integrated Project Management is to establish and manage the project and the involvement of the relevant stakeholders according to an integrated and defined process that is tailored from the organization's set of standard processes.

SG 1 Use the Project's Defined Process
The project is conducted using a defined process that is tailored from the organization's of standard processes.

SG 2 Coordinate and Collaborate with Relevant Stakeholders
The project coordinates and collaborates with the relevant stakeholders.

GG1-3

Risk Management – RM

Purpose

The purpose of Risk Management is to identify potential problems before they occur, so that risk-handling activities may be planned and invoked as needed across the life cycle to mitigate adverse impacts on achieving objectives.

SG 1 Prepare for Risk Management
Preparation for risk management is conducted.

SG 2 Identify and Analyze Risks
Risks are identified and analyzed to determine their relative importance.

SG 3 Mitigate Risks
Risks are handled and mitigated, where appropriate, to reduce adverse impacts on achieving objectives.

GG1-3

Decision Analysis and Resolution – DAR

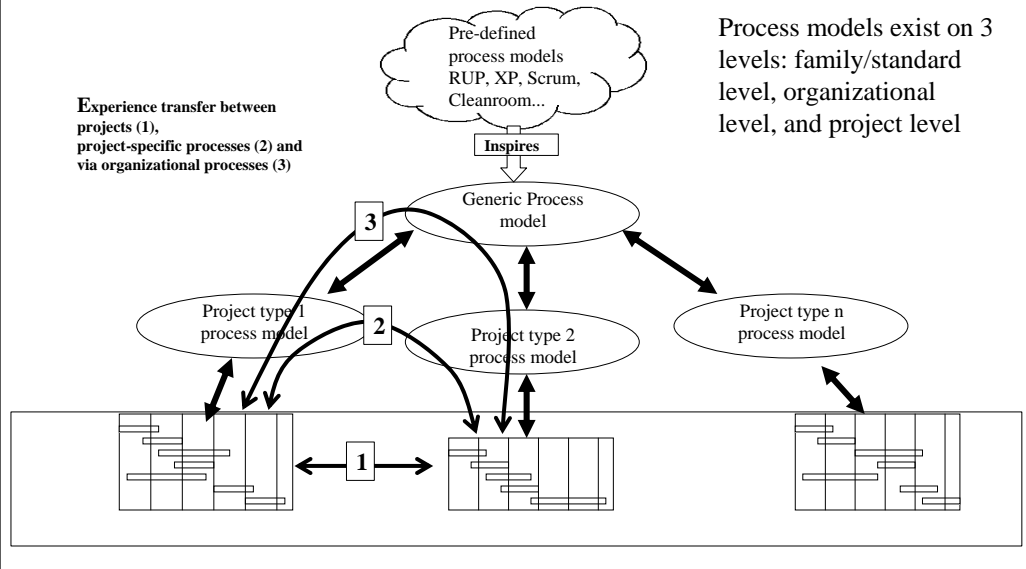
Purpose

The purpose of Decision Analysis and Resolution is to make decisions using a structured approach that evaluates identified alternatives against established criteria.

SG 1 Evaluate Alternatives
Decisions are based on an evaluation of alternatives using established criteria.

GG1-3

Learning and Improvement in SW Organisations



EVALUATION



CMMI Evaluation – How to do it?

Many models, forms and formalisms exist:

- Official *appraisal*: SCAMPISM (replaced CBA IPI* and SCE**)
- Company-specific assessments (e.g., Siemens Assessment)
- Light assessments
- Ultra-light assessments
- Self-assessment
- Interim-evaluation / Mini-assessment
- Based on open interviews
- Based on structured interviews
- Based on questionnaires

* CMM-Based Appraisal for Internal Process Improvement
** Software Capability Evaluation



CMMI Evaluation – Questionnaire Example

- **Related to Requirements Development**
- Note: This example refers to the SPICE model (cf. lecture Part 10), process ENG.2.2 - Analyze Software Requirements
- Similar questionnaires exist for CMMI (and other process assessment approaches)
- These questionnaires are NOT standardised

REQUIREMENTS ANALYSIS			
• Functional activities			
> Project development			
1. Was more detailed information about the software requirements gathered from the customer requirements document, internal constraints and other sources supplied by the customer?			
Observations: <input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
• Is it documented?			
<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
• Was it carried out according to an established procedure?			
<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
• Is the procedure documented?			
<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
2. Was a logic model of the product built from the functional requirements and the development methodology selected for the project?			
Observations: <input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
• Is it documented?			
<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
• Was it carried out according to an established procedure?			
<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
• Is the procedure documented?			
<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Not applicable <input type="checkbox"/> Don't know			
3. Were the requirements obtained from the customer analysed and verified to check that they were satisfactory and to detect any errors with regard to ambiguous, incomplete, unfeasible requirements, etc.?			
4. Was the logic model of the product validated against the functional requirements to assure that it satisfactorily met customer specifications?			
5. Was a prioritized list of requirements drawn up to enable gradual product development?			
6. Was a software requirements document drawn up on the basis of customer requirements internal organizational constraints?			
7. Was the software requirements document revised and approved by the customer?			
7.1. Was the software requirements document presented formally and comprehensibly to the customer?			
7.2. In the event that modifications were proposed to the software requirements document, Was their impact on the product logic model assessed?			
7.3. Did the customer approve the software requirements document in writing?			
8. Was the software requirements document managed and controlled?			
Management and control imply that a check is kept on versions and changes to the Software Requirements Document, that is, that the latest version is known at all times and amendments are introduced in a controlled manner.			
9. Was the software requirements document delivered to the Software Configuration Management group after acceptance and approval?			



SCAMPISM

- **Standard CMMI Appraisal Method for Process Improvement**

- Is a group of evaluation methods, both suitable to monitor progress on the way and for reliable benchmarking of organizations
- Complies with ISO 15504



The SCAMPISM Family

	Name:	Class A¹	Class B²	Class C³
	Type:	Benchmark	Mini-appraisal	Pulse Taking
	Objectiveness:	High	Medium	Low
	Evidence required:	Document Review Interviews Instrument	Document Review Interviews Instrument <i>(pick two)</i>	Document Review Interviews Instrument <i>(pick one)</i>
	Rating:	Formal Rating	Not formal	Not formal
	Responsible:	Lead Appraiser	Lead Appraiser <i>(preferred)</i>	Trained Leader
	Team:	Appraisal Team <i>(large)</i>	Appraisal Team <i>(medium)</i>	Appraisal Team <i>(small)</i>



Appraisal Team Members (ATM) – Requirements

- Appraisal Team must
 - have a total of 25 years of field experience with an average of 6 years
 - have a total of 10 years of management experience and at least one ATM with at least 6 years experience as a manager
 - have experience in the life cycles in use by the organizational unit
 - At least two ATMs should have experience as practitioners
- ATMs should
 - not be managers of any of the selected projects
 - not be in the direct supervisory chain of any of the interviewees
 - have good written and oral communication skills
 - have the ability to facilitate the free flow of communication
 - have the ability to perform as team players and negotiate consensus
 - have participated (at least 50%) in a previous process appraisal
 - be perceived by the appraisal sponsor as credible

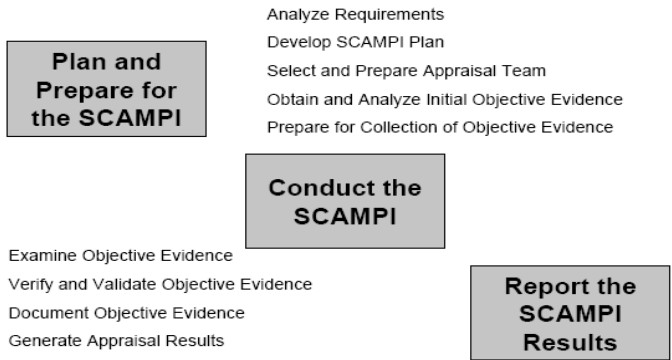


Practice Implementation Indicator Descriptions

PIID Type	Description	Examples
Direct	Tangible output resulting directly from implementation of a practice	Typical work products
Indirect	Artifacts that are a side-effect or indicative of performing a practice	Typical work products, meeting minutes, reviews, logs, reports
Affirmation	Oral or written statements confirming or supporting implementation of the practice	Interviews, questionnaires, briefings, demonstrations



The SCAMPI Process



TeraQuest



The SCAMPI Process (cont'd)

Phase	Process	Purpose	Activities
1 Plan and Prepare for Appraisal	1.1 Analyze Requirements	Understand the business needs of the organizational unit for which the appraisal is being requested. The appraisal team leader will collect information and help the appraisal sponsor match appraisal objectives with their business objectives.	1.1.1 Determine Appraisal Objectives 1.1.2 Determine Appraisal Constraints 1.1.3 Determine Appraisal Scope 1.1.4 Determine Outputs 1.1.5 Obtain Commitment to Appraisal Input
	1.2 Develop Appraisal Plan	Document requirements, agreements, estimates, risks, method tailoring, and practice considerations (e.g., schedules, logistics, and contextual information about the organization) associated with the appraisal. Obtain, record, and make visible the sponsor's approval of the appraisal plan.	1.2.1 Tailor Method 1.2.2 Identify Needed Resources 1.2.3 Determine Cost and Schedule 1.2.4 Plan and Manage Logistics 1.2.5 Document and Manage Risks 1.2.6 Obtain Commitment to Appraisal Plan
	1.3 Select and Prepare Team	Ensure that an experienced, trained, appropriately qualified team is available and prepared to execute the appraisal process.	1.3.1 Identify Team Leader 1.3.2 Select Team Members 1.3.3 Prepare Team
	1.4 Obtain and Analyze Initial Objective Evidence	Obtain information that facilitates site-specific preparation. Obtain data on model practices used. Identify potential issue areas, gaps, or risks to aid in refining the plan. Get preliminary understanding of the organizational unit's operations and processes.	1.4.1 Prepare Participants 1.4.2 Administer Instruments 1.4.3 Obtain Initial Objective Evidence 1.4.4 Inventory Objective Evidence
	1.5 Prepare for Collection of Objective Evidence	Plan and document specific data collection strategies including sources of data, tools and technologies to be used, and contingencies to manage risk of insufficient data.	1.5.1 Perform Readiness Review 1.5.2 Prepare Data Collection Plan 1.5.3 Replan Data Collection (if needed)



The SCAMPI Process (cont'd)

Phase	Process	Purpose	Activities
2 Conduct Appraisal	2.1 Examine Objective Evidence	Collect information about the practices implemented in the organizational unit and relate the resultant data to the reference model. Perform the activity in accordance with the data collection plan. Take corrective actions and revise the data collection plan as needed.	2.1.1 Examine Objective Evidence from Instruments 2.1.2 Examine Objective Evidence from Presentations 2.1.3 Examine Objective Evidence from Documents 2.1.4 Examine Objective Evidence from Interviews
	2.2 Verify and Validate Objective Evidence	Verify the implementation of the organizational unit's practices for each instantiation. Validate the preliminary findings, describing gaps in the implementation of model practices. Each implementation of each practice is verified so it may be compared to CMMI practices, and the team characterizes the extent to which the practices in the model are implemented. Gaps in practice implementation are captured and validated with members of the organizational unit. Exemplary implementations of model practices may be highlighted as strengths to be included in appraisal outputs.	2.2.1 Verify Objective Evidence 2.2.2 Characterize Implementation of Model Practices 2.2.3 Validate Practice Implementation Gaps
	2.3 Document Objective Evidence	Create lasting records of the information gathered by identifying and then consolidating notes, transforming the data into records that document practice implementation, as well as strengths and weaknesses.	2.3.1 Take/Review/Tag Notes 2.3.2 Record Presence/Absence of Objective Evidence 2.3.3 Document Practice Implementation Gaps 2.3.4 Review and Update the Data Collection Plan
	2.4 Generate Appraisal Results	Rate goal satisfaction based upon the extent of practice implementation throughout the organizational unit. The extent of practice implementation is determined/judged based on validated data (e.g., the three types of objective evidence) collected from the entire representative sample of the organizational unit. The rating of capability levels and/or maturity levels is driven algorithmically by the goal satisfaction ratings.	2.4.1 Derive Findings and Rate Goals 2.4.2a Determine Process Area Capability Level 2.4.2b Determine Satisfaction of Process Areas 2.4.3a Determine Capability Profile 2.4.3b Determine Maturity Level 2.4.4 Document Appraisal Results

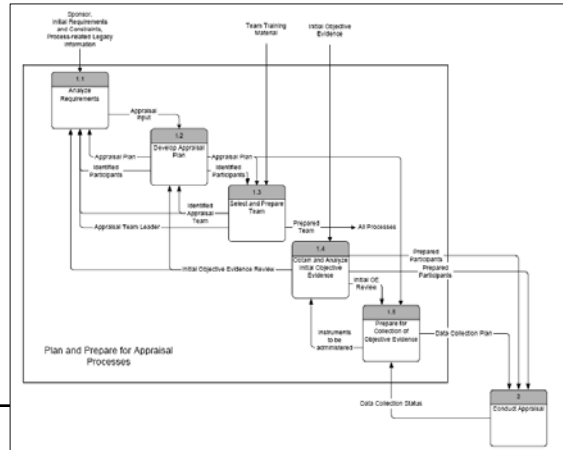


The SCAMPI Process (cont'd)

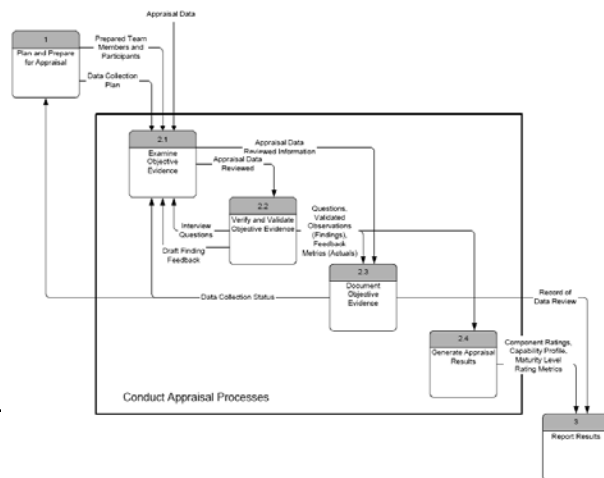
Phase	Process	Purpose	Activities
3 Report Results	3.1 Deliver Appraisal Results	Provide credible appraisal results that can be used to guide actions. Represent the strengths and weaknesses of the processes in use at the time. Provide ratings (if planned for) that accurately reflect the capability level/maturity level of the processes in use.	3.1.1 Present Final Findings 3.1.2 Conduct Executive Session(s) 3.1.3 Plan for Next Steps
	3.2 Package and Archive Appraisal Assets	Preserve important data and records from the appraisal, and dispose of sensitive materials in an appropriate manner.	3.2.1 Collect Lessons Learned 3.2.2 Generate Appraisal Record 3.2.3 Provide Appraisal Feedback to CMMI Steward 3.2.4 Archive and/or Dispose of Key Artifacts



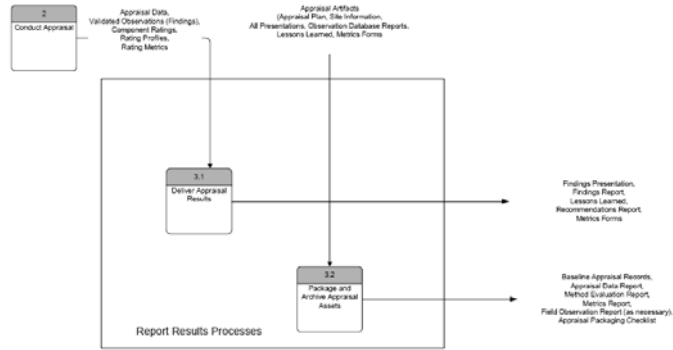
The SCAMPI Process – Control Flow



The SCAMPI Process – Control Flow (cont'd)

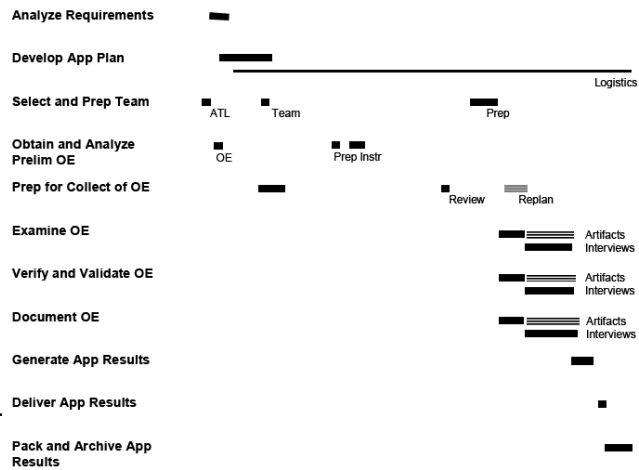


The SCAMPI Process – Control Flow (cont'd)



SCAMPI – Nominal Assessment Schedule View

ATL = Appraisal Team Leader
 OE = Objective Evidence



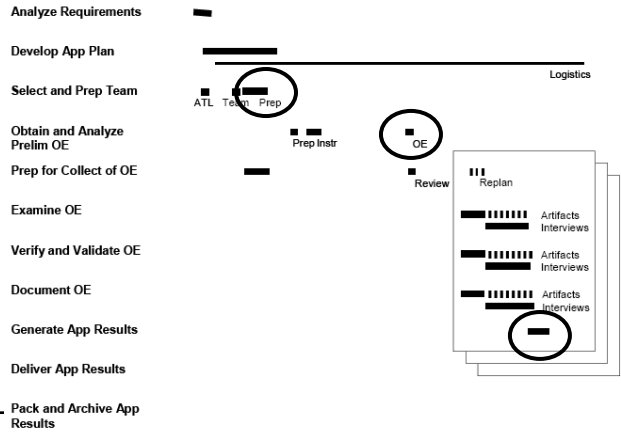
SCAMPI – Nominal Evaluation Schedule View

The basic difference between an assessment and an evaluation is that an assessment is an appraisal that an organization does to and for itself for the purposes of process improvement.

Assessments provide internal motivation for organizations to initiate or continue process improvement programs.

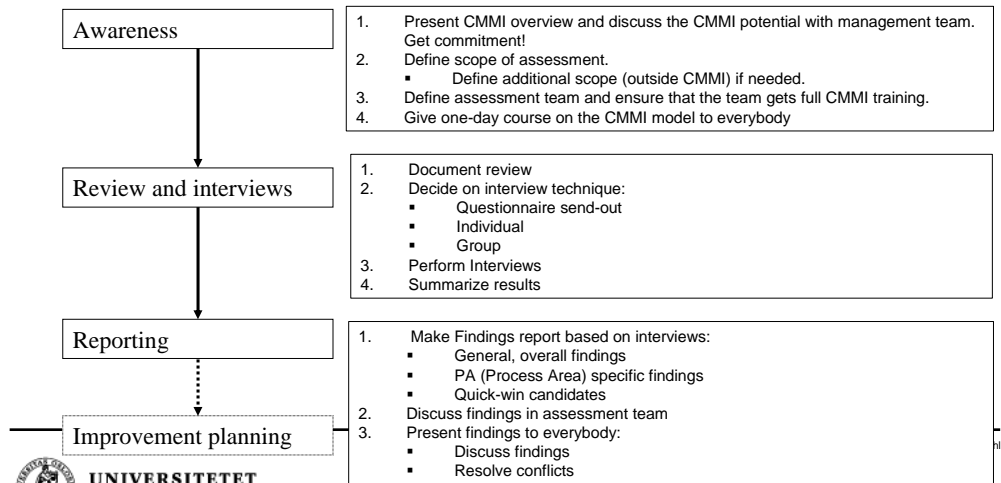
An evaluation is an appraisal in which an external group comes into an organization and examines its processes as input to a decision regarding future business.

Evaluations are typically externally imposed motivation for organizations to undertake process improvement.



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A Simple Assessment-Process



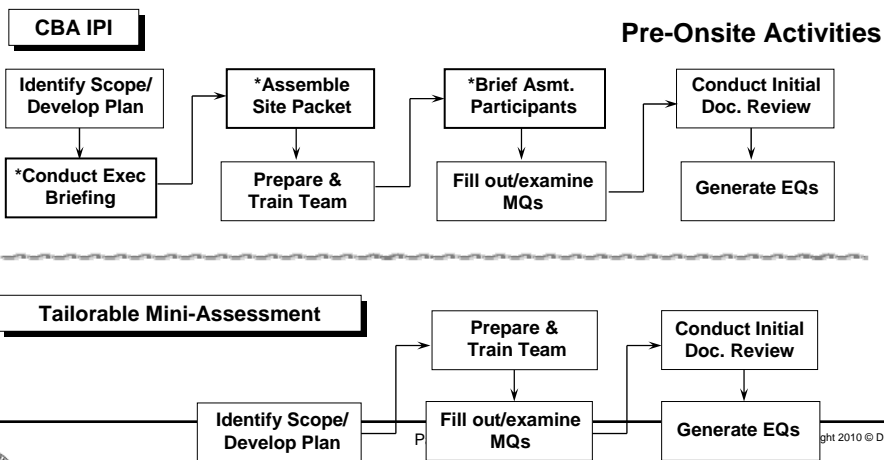
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Mini-Assessment Method

- An informal review of an organization's current software process based on:
 - A review of 3-4 key projects
 - responses to SEI's MQ (Maturity Questionnaire)
 - discussions with senior managers, project leaders, middle managers, and practitioners
 - document review
- Uses a tailored and streamlined version of the SEI's CBA IPI method
 - Cost and resource impacts reduced
 - less time, fewer participants, some sessions combined or deleted, less formality
 - No ratings
 - Focuses on global strengths and high priority issues
 - primarily weaknesses at the KPA level



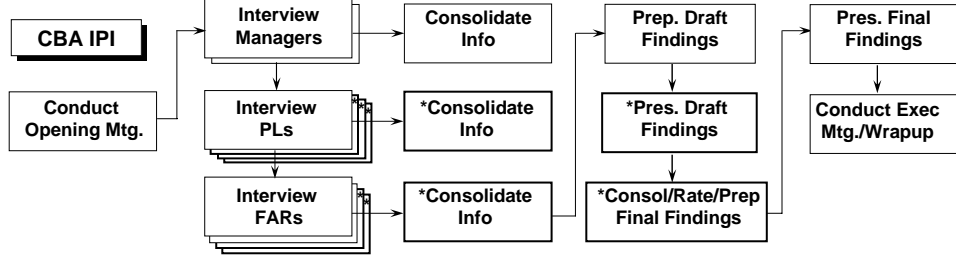
CBA IPI & Mini-Assessment Comparison



*Asterisked (red items) are omitted during a mini-assessment.
EQs=Exploratory Questions MQs=Maturity Questionnaires

CBA IPI & Mini-Assessment Comparison

Onsite Activities



Notes: * Asteriskd (red) items are omitted during a mini-assessment. Managers=Senior & Middle Mgrs. PLs=Project Leads FARs=Functional Area Reps



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Page 73

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Mini-Assessment Status Chart (optional)

SOFTWARE PROJECT PLANNING								
	Commitment to Perform	Ability to Perform	Activities Performed				Measurement & Analysis	Verifying Implementation
			1	2	3	4		
Common Feature Definitions:								
Commitment to Perform:	documented policy, senior management commitment demonstrated, responsibilities assigned							
Ability to Perform:	adequate resources (tools, staff, \$) to perform the practices, role/function in place, trained personnel							
Activities Performed:	1. processes/template documented & used 2. estimates, commitments, & schedules generated/documentd							
Measurement & Analysis:	3. project plans complete & documented, 4. risks (technical & program) identified							
Verifying Implementation:	process metrics collected & analyzed							
Compliance Legend:	process activities reviewed by senior management, middle management, & SQA							
	Practices implemented and institutionalized (in place on all assessed projects)							
	Some practices implemented. Or practices performed only some projects (inconsistent across the organization).							
	Little or no evidence of practices in place (on the assessed projects).							

Page 74

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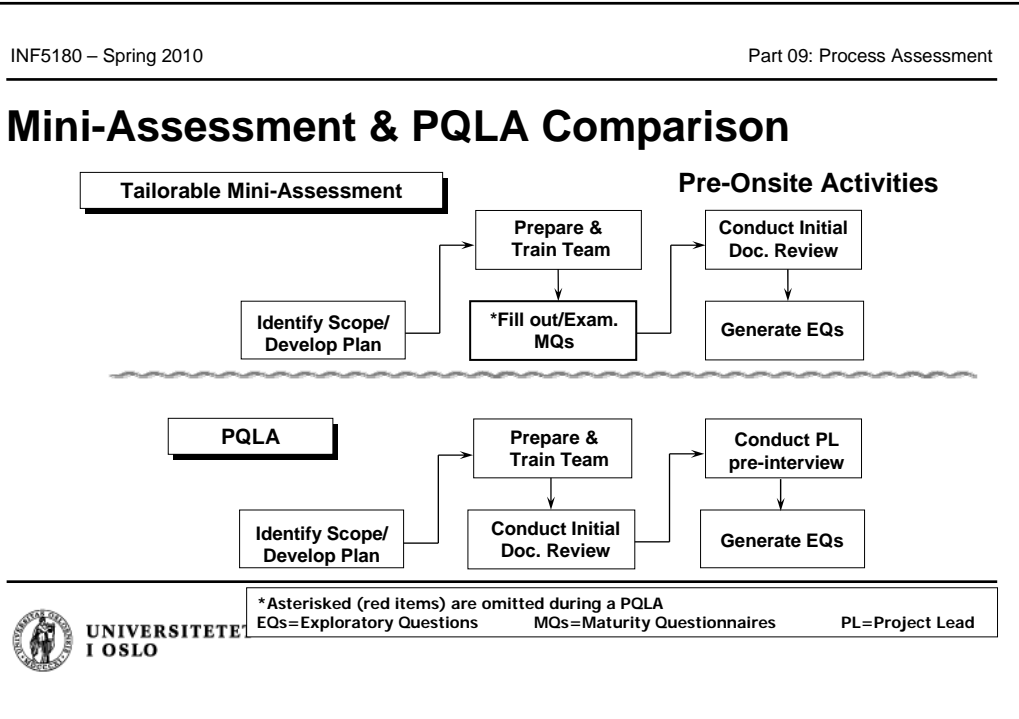


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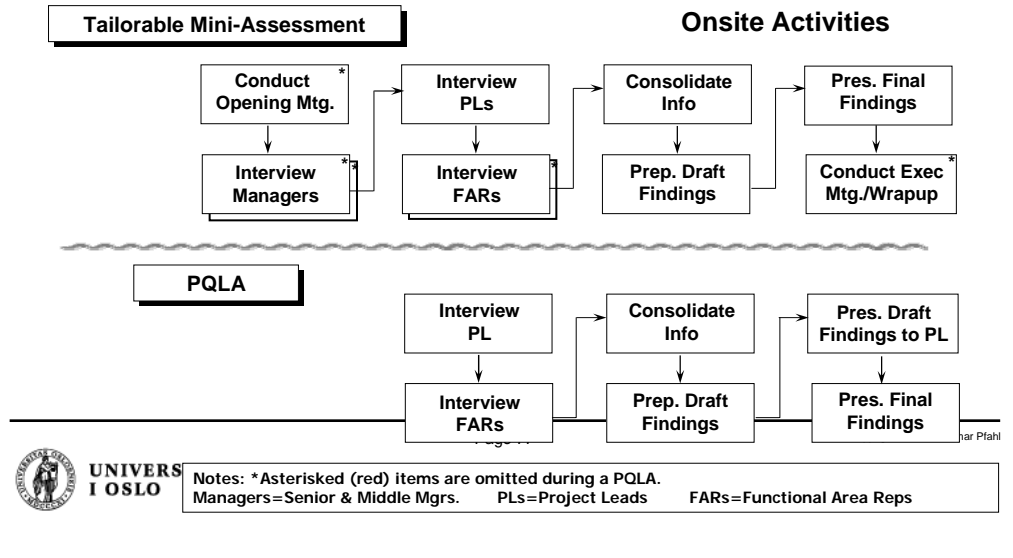
Project Quick Look Assessment Method

- An informal review of a single project’s current software process based on:
 - Discussions with Project Lead and practitioners
 - Document review
- Uses a tailored and streamlined version of the mini-assessment method
 - Cost and resource impacts greatly reduced
 - minimal time, fewer participants, some sessions combined or deleted, informal discussions and briefing
 - Reduced scope
 - single project only; no organizational practices evaluated
 - Interview questions more direct and interactive
 - No ratings
 - Focuses on high priority issues at the KPA level

PQLA approach developed and successfully used at General Dynamics

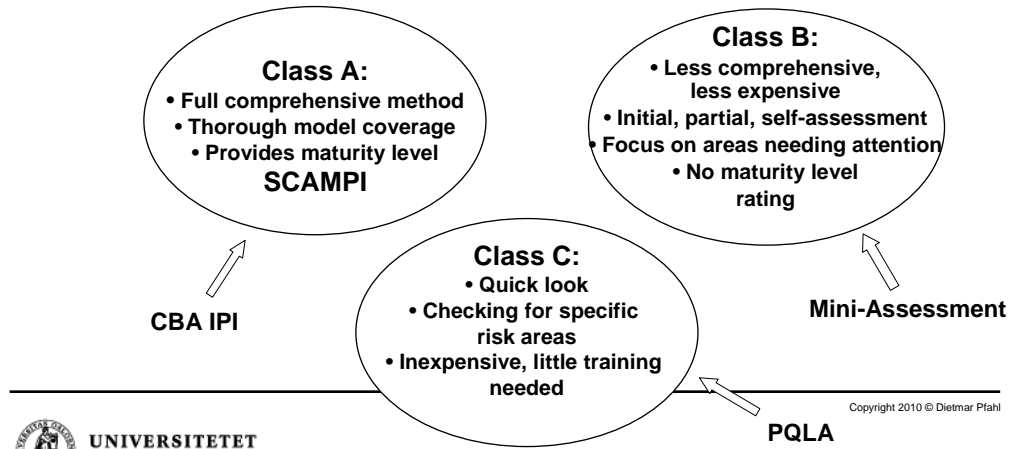


Mini-Assessment & PQLA Comparison



CMMIsm Classes of Assessment Methods

Comparison with CMM Assessment Methods



SCAMPI = Standard CMMI Assessment Method for Internal Process Improvement.
 sm CMM Integration and CMMI are service marks of Carnegie Mellon University.

Assessment Method Comparisons

	CBA IPI	Mini-asmt.	PQLA
*Resources:			
- # team members	6-8	4-6	4
- team member time (plan, prep, conduct)	110-130 hrs.	48-60 hrs.	14-20 hrs.
- # participants	50-60	30-40	8-10
- participant time (prep, conduct)	4-8 hrs.	2-5 hrs.	1-3 hrs.
Team training (CMM and assessment method)	5 days	1.5-2 days	4-6 hrs.
Pre On-Site schedule (wall time)	2-3 months	3-4 weeks	1 week
On-Site schedule (consecutive days)	7-9 days	4-5 days	1.5-2 days
Formality (briefings, plans, reports, paperwork)	<ul style="list-style-type: none"> • Formal • Maximum doc. review 	<ul style="list-style-type: none"> • Informal • Moderate doc. review 	<ul style="list-style-type: none"> • Very informal • Minimal doc. review

*Resources are per person; Typical figures for an organization with size 100 SW staff, covering Levels 2 & 3. Total time includes planning, preparing, and conducting.



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Assessment Method Comparisons (cont.)

	CBA IPI	Mini-assessment	PQLA
Outputs	<ul style="list-style-type: none"> • Findings briefing: <ul style="list-style-type: none"> - Global findings - KPA findings (strengths & weaknesses) - Maturity Level - KPA ratings • Final Report • Data/results to SEI 	<ul style="list-style-type: none"> • Findings briefing: <ul style="list-style-type: none"> - Global findings - KPA findings (strengths & weaknesses) • Color chart (opt) 	<ul style="list-style-type: none"> • Findings briefing: <ul style="list-style-type: none"> - Global findings - KPA weaknesses
Pros	<ul style="list-style-type: none"> • Very comprehensive / accurate • Supports detailed action plan 	<ul style="list-style-type: none"> • Comprehensive • Reliable predictor of CBA IPI results • Less time, \$, participants, tension 	<ul style="list-style-type: none"> • Minimal time, \$, participants • Participants more at ease; interactive
Cons	<ul style="list-style-type: none"> • Expensive • Time consuming • Schedule difficulties • Tension due to ratings 	<ul style="list-style-type: none"> • Schedule difficulties 	<ul style="list-style-type: none"> • Some weaknesses may be missed • Does not provide organizational view



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Interim Evaluation – CMM-Mini

- A light weight evaluation for projects and line organization – developed by KDA
- Tool: a spreadsheet that lists all relevant KPAs with its practices
- All practices are given a score:
 - y – Yes
 - p – Partly
 - n – Not
 - na – Not Applicable
- Assumes that mapping to procedures and documents (PIIDs) has been done in advance.
- Project:
 - All KPAs and all practices which are relevant for the project (all in level 2 plus a selection in level 3)
 - Are implemented regularly (every or every other month)
 - Are based on 1 full review (< 8 hours), while the other is focused (< 2 hours)
- Line: A selection of KPAs on level 3 where the responsibility is centralized.



Example: Requirement Management

Requirements Management

Goals for the RM Key Process Area:

Goal 1: System requirements allocated to software are controlled to establish a baseline for software engineering and management use.

Goal 2: Software plans, products and activities are kept consistent with the system requirements allocated to software.

Key practices	Values	Assess the status in the "Values" column; y=yes, p=partly, n=no	How is it handled within the project?	Solutions provided from the line organization	Action required to achieve compliance?
c	1	y	The project follows a written organizational policy for managing the allocated system requirements.		KDA Quality Manual, part I SE-process and INS 0234
ab	1	y	For each project, responsibility is established for analysing the system requirements and allocating them to HW, SW, and other system components	Responsibility (Role) defined and allocated to a person or group.	KDA Roles: SE-manager, responsible for allocating requirements to SW, HW etc. SW architect responsible for writing the SRS.
ab	2	y	The allocated requirements are documented.	Technical, Non-technical requirements and acceptance criteria are defined and documented!	SE-process: User Requirement specification (URS), System Segment Specification (SSS), Interface requirement specification (IRS), Software Requirement Specification (SRS), Hardware development specification (HDS), Statement of Work (SOW) INS 0234
ab	3	y	Adequate resources and funding are provided for managing the allocated requirements.	An agreement written between the line and project for the allocated RM resources. Hours for RM planned, Necessary tools available.	AZDS tool: Rational Requisite Pro
ab	4	y	Members of the engineering group and other related groups are trained to perform their requirements management activities.	Training in RM activities both technical application, methods and tools. Example Prephase training.	
ac	1	y	The engineering group reviews the allocated requirements before they are incorporated into the project.		PRO-0017 Review Process, SRS checklist in PRO-0016, AZDS Requirements to projects: Requirements handling.
ac	2	y	The engineering group uses the allocated requirements as the basis for plans, work products, and activities.		Project management process, Initial planning PRO-0015 Software Engineering Management, Plan the SW development uses SRSs as input to planning. AZDS Task descriptions?



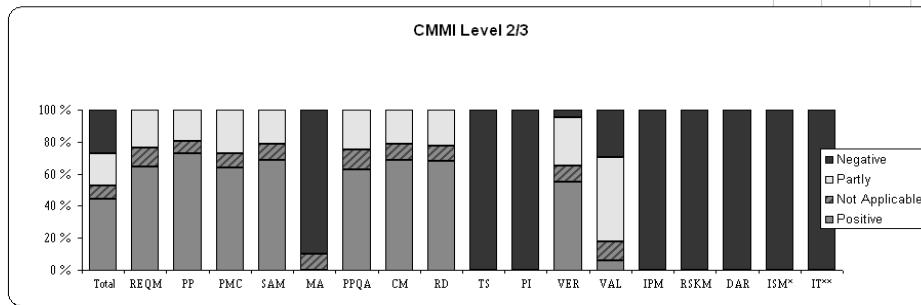
CMM-Mini Results (project)

Total CMMI Compliance: 63 %

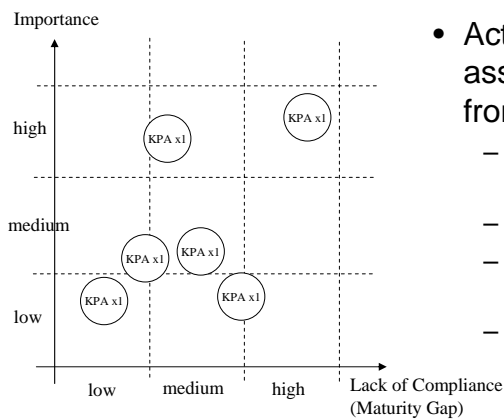
Tailoring: Use of Not Applicable: 8 %

CMMI Level 2 compliance: 77 %

CMMI Level 3 compliance: 22 %



Assessment Follow-On Activities



- Action plan – generated by SEPG, assessment team, and key personnel from organization/projects
 - Address findings (how to address weaknesses)
 - Strategy for addressing additional KPAs
 - Detailed actions, responsibilities, budget, and schedule
 - Reviewed/approved by management





and ...

CONTINUOUS PROCESS IMPROVEMENT



SEI Slogan

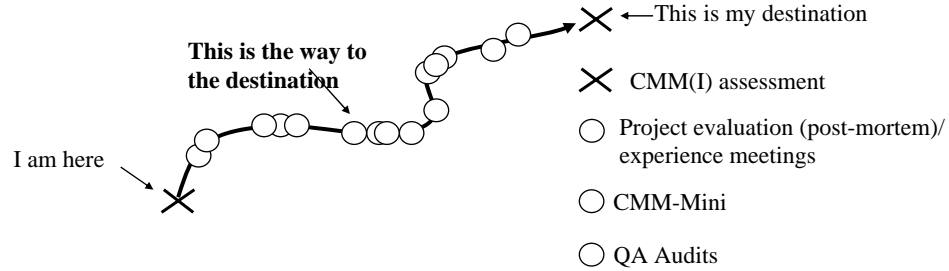
“The Capability Maturity Model for Software (CMM) is a framework that describes the elements of an effective software process.

The CMM describes an evolutionary path from an ad hoc, chaotic process, to a mature disciplined process”



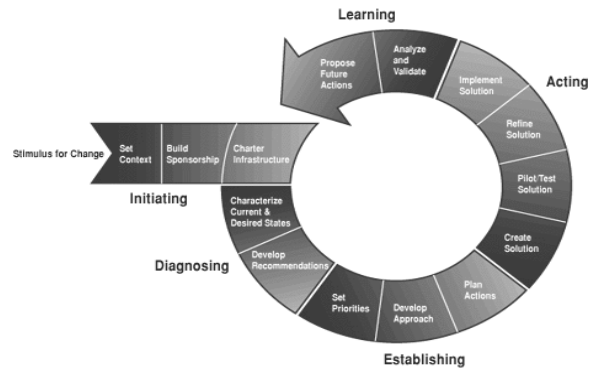
CMMI as a Roadmap

- It is not difficult to come up with a long list with good proposals for improvements.
- It is more difficult to prioritize the most important and to make a realistic progress plan.
- The most difficult of all is to manage the necessary changes in the organization – work pattern and not the least culture.



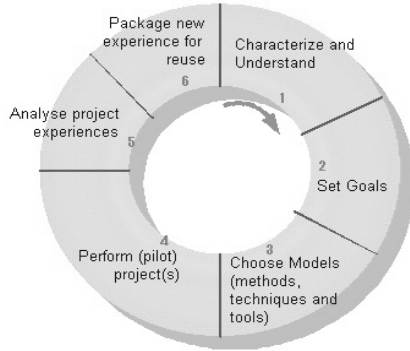
Process Improvement: IDEAL

The Process Improvement Process of SEI:



CMM and QIP

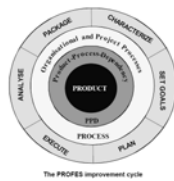
- CMM is well suited to be used with Quality Improvement Paradigm on the organizational level



1. Conduct a CMM-assessment. Combine the results with business goals and identified problems.
2. Use "Findings" from the assessment to define improvement goals
3. Choose measures which are supposed to help achieve the objectives
4. Implement one or more projects with the new models, tools and methods. Collect data on the way
5. Collect lessons learnt from the projects
6. Update standard process model and experience database



PROFES – Product-Focused Process Improvement in Software Engineering



PROFES PHASES	PROFES STEPS
CHARACTERIZE	1. VERIFY COMMITMENT
	2. IDENTIFY PRODUCT QUALITY NEEDS
	3. DETERMINE CURRENT PRODUCT QUALITY
	4. DETERMINE CURRENT PROCESS CAPABILITY
SET GOALS	5. SET PRODUCT IMPROVEMENT GOALS
	6. DETERMINE NECESSARY PROCESS CHANGES
PLAN	7. DESCRIBE PROCESS CHANGES
	8. SET METRICS FOR THE PROCESSES AND PRODUCT
	9. PREPARE IMPROVEMENT IMPLEMENTATION
EXECUTE	10. IMPLEMENT AND MONITOR IMPROVEMENTS
ANALYSE	11. EVALUATE RESULTS
PACKAGE	12. UPDATE EXPERIENCE BASE

Phases and steps of the PROFES improvement methodology

Step 1 Verify commitment

Goals	Activities
<ul style="list-style-type: none"> The organization's business needs and improvement objectives for product and process quality are identified. Product quality characteristics, ongoing improvement initiatives, and their priorities are identified. Commitment of top and middle management is verified. Commitment of project members is verified. Current state info of the organization & projects is defined. An overall plan for improvement activities is defined. 	<ul style="list-style-type: none"> Identify the organization's business needs and improvement objectives Motivate top and middle management Mobilize project members Define organizational context Define overall plan and schedule
Input	Output
<ul style="list-style-type: none"> Organizational level: <ul style="list-style-type: none"> General organizational information Business goals Customer survey results Market research results Customer feedback Organizational context information Project level: <ul style="list-style-type: none"> Project environment specifics Product development goals 	<ul style="list-style-type: none"> Organizational level: <ul style="list-style-type: none"> Commitment of top and middle management Preliminary product and process improvement needs Organization's classification Overall improvement plan Project level: <ul style="list-style-type: none"> Commitment of project management and members Project classification Overall improvement plan

Step 2 Identify product quality needs

Goals	Activities
<ul style="list-style-type: none"> Product quality needs are known and presented in the form of a product quality profile Preliminary product quality goals are set 	<ul style="list-style-type: none"> Survey product quality needs Document product quality needs Set preliminary product quality goals
Input	Output
<ul style="list-style-type: none"> Customer survey results Market research results Customer feedback Business goals IS/IMS Preliminary product quality needs 	<ul style="list-style-type: none"> Product quality needs Product quality profile Preliminary product quality goals

Step 3 Determine current product quality

Goals	Activities
<ul style="list-style-type: none"> Determine current status of product quality 	<ul style="list-style-type: none"> Acquire product quality data Evaluate current status of product quality
Input	Output
<ul style="list-style-type: none"> Applicable domain characteristics Measurement data IS/IMS Product quality profile Experience base 	<ul style="list-style-type: none"> Current status of product quality

Step 4 Determine current process capability

Goals	Activities
<ul style="list-style-type: none"> Current process capability is determined Process improvement recommendations are documented and communicated 	<ul style="list-style-type: none"> Preparation Execution Reporting
Input	Output
<ul style="list-style-type: none"> Business goals Process descriptions Quality manuals Organizational characteristics Project plans Design documents Measurement data 	<ul style="list-style-type: none"> Process capability profiles Process assessment report and profiles Descriptive process models Preliminary improvement plan



<p>Step 5 Set product improvement goals</p> <p>Goals</p> <ul style="list-style-type: none"> Set Product improvement goals <p>Activities</p> <ul style="list-style-type: none"> Analyse product quality discrepancies Identify product improvement areas Prioritize product improvement areas Set the product improvement goals <p>Output</p> <ul style="list-style-type: none"> Product improvement goals <p>Business goals</p> <p>Product quality needs</p> <ul style="list-style-type: none"> Product quality target profile Current status of product quality Process assessment reports and profiles Descriptive process models Preliminary product quality goals Product characteristics 	<p>Step 9 Prepare improvement implementation</p> <p>Goals</p> <ul style="list-style-type: none"> Plan process changes and allocate sufficient resources to implement them Plan improvement progress meetings <p>Activities</p> <ul style="list-style-type: none"> Plan process improvement progress meetings Make time planning and resource allocation Kick-off process changes <p>Input</p> <ul style="list-style-type: none"> Development project plan Preliminary improvement plan (from Step 4) Selected list of process changes (from Step 6) Prescriptive process model (from Step 7) GQM deliverables (from Step 8) <p>Output</p> <ul style="list-style-type: none"> Process improvement action plan On-line process support
<p>Step 6 Determine necessary process changes</p> <p>Goals</p> <ul style="list-style-type: none"> Identify and select process changes necessary to achieve the product improvement goals Document the decisions on necessary process changes for later evaluation of the improvement programme <p>Activities</p> <ul style="list-style-type: none"> Identify product quality goal Identify processes to be improved Retrieve relevant PPD models Construct characterization questionnaire Characterize the project Rank PPD models Select improvement actions <p>Output</p> <ul style="list-style-type: none"> Process changes to be implemented in the improvement programme Characterization of the forthcoming project or improvement programme <p>Input</p> <ul style="list-style-type: none"> Product improvement goals Process assessment reports and profiles (from Step 4) PPD repository Preliminary improvement plan (from Step 4) 	<p>Step 10 Implement and monitor improvements in the development project</p> <p>Goals</p> <ul style="list-style-type: none"> Implement selected process changes according to process improvement plan Collect data and prepare measurement results for each feedback session Meet feedback sessions <p>Activities</p> <ul style="list-style-type: none"> Implement process changes Collect measurement data Prepare and select measurement data Perform GQM feedback sessions <p>Input</p> <ul style="list-style-type: none"> Prescriptive process model GQM plan Measurement plan Process improvement plan Development project plan <p>Output</p> <ul style="list-style-type: none"> Measurement data Feedback session report(s) with visualized measurement data Description of corrective actions taken Prescriptive process model applied in practice
<p>Step 7 Describe process changes</p> <p>Goals</p> <ul style="list-style-type: none"> Agree and document prescriptive process model Achieve clear understanding of the processes in order to define the metrics in the following step <p>Activities</p> <ul style="list-style-type: none"> Mark processes/practices in the current process model, which have to be changed Develop prescriptive process model Communicate prescriptive model to process participants <p>Output</p> <ul style="list-style-type: none"> Prescriptive process model (including selected process changes) Training/presentation material for the new process <p>Input</p> <ul style="list-style-type: none"> Descriptive process model (from Step 4) Selected list of process changes (from Step 6) 	<p>Step 11 Evaluate Results</p> <p>Goals</p> <ul style="list-style-type: none"> Evaluate effect of the improvement programme on final product quality Evaluate changes to the software engineering process, methods, and tools Gather and evaluate "lessons learned" Support, modify, or reject used PPD models <p>Activities</p> <ul style="list-style-type: none"> Evaluate the measurement results Support, modify, or reject used PPD models <p>Input</p> <ul style="list-style-type: none"> PPD models Prescriptive process model Abstraction sheets GQM plan GQM measurement plan Measurement data Feedback session reports <p>Output</p> <ul style="list-style-type: none"> Preliminary experience packages Evaluated PPD models
<p>Step 8 Set metrics for the processes and product improvements</p> <p>Goals</p> <ul style="list-style-type: none"> Define questions and metrics related to the product quality goals Define questions and metrics related to the process performance goals Define questions and metrics related to the product-process dependency goals Construct GQM plan and measurement plan <p>Activities</p> <ul style="list-style-type: none"> Define measurement goals Conduct GQM interviews Define questions and hypotheses Define and check metrics Prototype GQM plan and measurement plan <p>Output</p> <ul style="list-style-type: none"> GQM abstraction sheets GQM plan Measurement plan <p>Input</p> <ul style="list-style-type: none"> Prescriptive process model (including selected process changes) Product quality and target profile (from Step 2) Current status of product quality (from Step 3) Product improvement goals (from Step 5) Process assessment reports and profiles (from Step 4) PPD models (from Step 6) 	<p>Step 12 Update Experience Base</p> <p>Goals</p> <ul style="list-style-type: none"> Package and store all information gained during the project in the experience base for future reuse <p>Activities</p> <ul style="list-style-type: none"> Package information Store relevant information in the experience base <p>Input</p> <ul style="list-style-type: none"> Evaluated PPD models Experience base Process models GQM plans Feedback session reports <p>Output</p> <ul style="list-style-type: none"> Updated experience base with generalized: <ul style="list-style-type: none"> PPD models Process models GQM plans

Different Process Improvement Approaches

- Most users of CMM(I) today use it internally as a improvement framework – i.e. they don't intend to use assessment results for marketing purpose.
- The most active CMM advocates (Mark Paulk and Bill Curtis) strongly warn against following the model blindly. *Use the model with common sense, coupled with understanding one's own needs and problem areas.*

Model-less SPI
(e.g. GQM)

CMM(I) used as guideline
(combined with GQM/QIP or similar)

"Hard-core CMM"
(SCAMPI)



Use Common Sense...

- Mark Paulk, SEI:

		Process Discipline	
		Yes	No
Common Sense	Yes	Quality	Creative Chaos
	No	Mindless Bureaucracy	Mindless Chaos



DISSEMINATION and RESULTS



Improvements According to SEI Data

Performance Results of CMMI®-Based Process Improvement

The quantitative performance results in Table 2 are from a total of 35 organizations, some of which are enterprises with more than one constituent organization. 30 of them have results that can be expressed as change over time. These results are expressed either as percentage change from an earlier baseline prior to the CMMI-based process improvement or as ratios of return on investment (ROI). The results are summarized by the six performance categories discussed in Section 2 of this document: cost, schedule, productivity, quality, customer satisfaction and return on investment. Most of the organizations have provided multiple results, sometimes several in the same performance category.

Table 2: CMMI Performance Results Summary

Performance Category	Median Improvement	Number of Data Points	Lowest Improvement	Highest Improvement
Cost	34%	29	3%	87%
Schedule	50%	22	2%	95%
Productivity	61%	20	11%	329%
Quality	48%	34	2%	132%
Customer Satisfaction	14%	7	-4%	55%
Return on Investment	4.0 : 1	22	1.7 : 1	27.7 : 1

Note: The performance results in this table express change over varying periods of time.

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CMU/SEI-2006-TR-004
ESC-TR-2006-004



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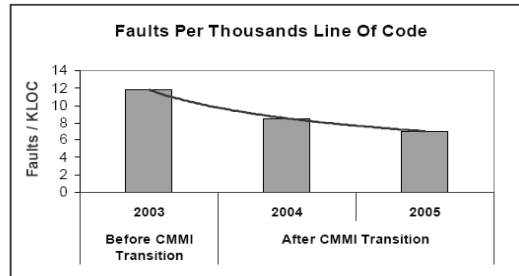
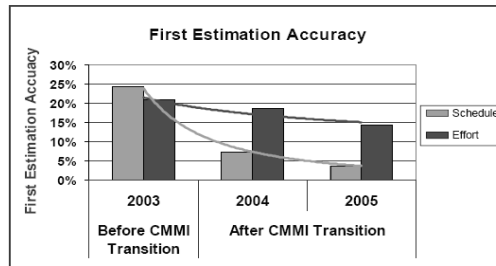


Figure 24: In Process Faults per Thousand Lines of Code



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Raytheon Network Centric Systems

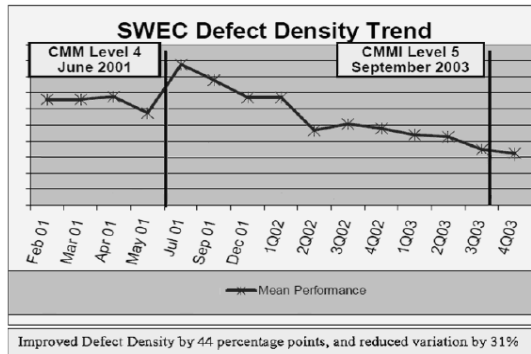
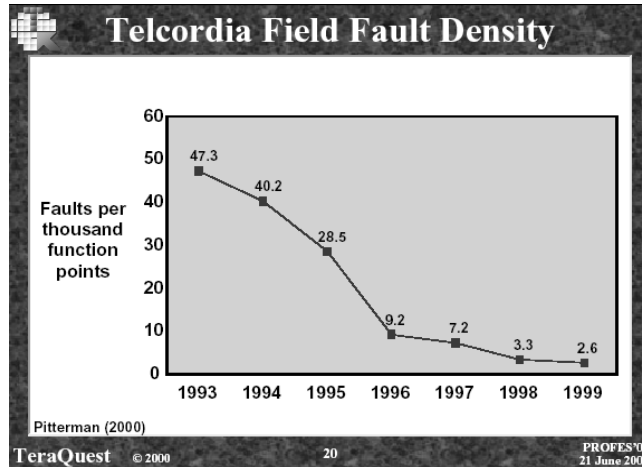


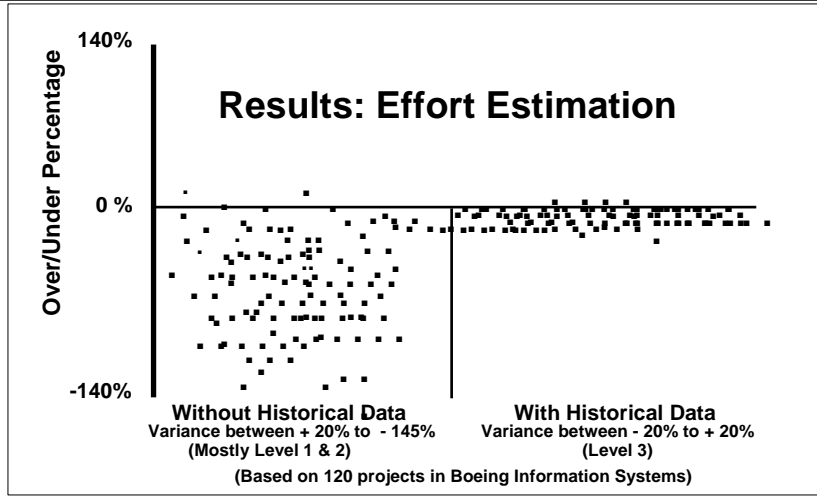
Figure 40: Defect Density



Telcordia (CMM)



Boeing



Reference: John D. Vu. "Software Process Improvement Journey: From Level 1 to Level 5." 7th SEPG Conference, San Jose, March 1997.

Status and Spread of CMMI

- SCAMPI v1.1 Class A appraisals conducted since its April 2002 release and reported to the SEI by July 2004
 - 367 appraisals
 - 333 organizations
 - 176 participating companies
 - 28 reappraised organizations
 - 1.368 projects
 - 46,2 % Non-USA organizations

Organization Type
Based on Primary Standard Industrial Classification (SIC) Code

