



Required, Expected, and Informative Components

- **Required**: Specific and generic goals (what must be implemented)
- Expected: Specific and generic practices (activities expected)
- Informative: Subpractices, typical work products, discipline amplifications, generic practice elaborations, goal and practice titles, goal and practice notes, and references

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3



Model Components -1

- Process areas: a PA is a cluster of related practices in an area, when performed
 collectively, satisfy a set of goals considered important for making significant
 improvement in that area.
- 2. Specific goals: apply to a process area and address the <u>unique characteristics</u> that describe what must be implemented to satisfy the PA.
- 3. Specific practices: a specific practice is an activity expected.
 - Base practices: level 1 practices
 - Advanced practices: levels of 2 or higher
- 4. Typical work products
 - Provide <u>example</u> outputs from a specific or generic practice.



Model Components -2

- 5. Subpractices: detailed descriptions that provide guidance for interpreting specific or generic practices.
- 6. Discipline amplification
- 7. Generic goals: describe the institutionalisation that the organization must achieve at that capability level.
- **8.** Generic practices: provide institutionalisation to ensure that the processes associated with the PA will be effective, repeatable, and lasting.
- 9. Generic practices elaborations
- 10. References

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Example: Requirements Development

SG 1 Develop Customer Requirements

SP 1.1-1 Collect Stakeholder Needs

SP 1.1-2 Elicit Needs

SP 1.2-1 Develop the Customer Requirements

SG 2 Develop Product Requirements

SP 2.1-1 Establish Product & Product-Component

SP 2.2-1 Allocate Product-Component Requirements

SP 2.3-1 Identify Interface Requirements

SG 3 Analyze and Validate Requirements

SP 3.1-1 Establish Operational Concepts and Scenarios

SP 3.2-1 Establish a Definition of Required Functionality

SP 3.3-1 Analyze Requirements

SP 3.4-3 Analyze Requirements to Achieve Balance

SP 3.5-1 Validate Requirements

SP 3.5-2 Validate Requirements with Comprehensive Methods

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

GG 4 Institutionalize a Quantitatively Managed

GP 4.1 Establish Quantitative Objectives for the Process

GP 4.2 Stabilize Subprocess Performance

GG 5 Institutionalize an Optimizing Process

GP 5.1 Ensure Continuous Process Improvement GP 5.2 Correct Root Causes of Problems

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25 Process Areas Continuous Representation

	Organizational Process Focus
Process management	Organizational Process Definition
	Organizational Training
	Organizational Process Performance
	Organizational Innovation and Deployment
1	Project Planning
	Project Monitoring and Control
Project management	Supplier Agreement Management
	Integrated Project Management for IPPD
	Risk Management
	Integrated Teaming (IPPD)
	Integrated Supplier Management (SS)
	Quantitative Project Management
	Requirements Management
	Requirements Development
	Technical Solution
Engineering	Product Integration
	Verification
	Validation
	Configuration Management
Support	Process and Product Quality Assurance
	Measurement and Analysis
	Decision Analysis and Resolution
	Organizational Environment for Integration (IPPD)
	Causal Analysis and Resolution

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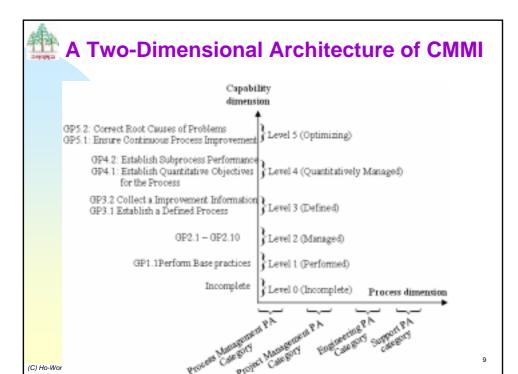
7



CMMI GPs

Capability level	Generic practices			
CL0 (Incomplete)				
CL1 (Performed)	GP1.1: Perform base practices			
CL 2 (Managed)	GP2.1: Establish an organizational policy			
	GP2.2: Plan the process			
	GP2.3: Provide resources			
	GP2.4: Assign responsibility			
	GP2.5: Train people			
	GP2.6: Manage configurations			
	GP2.7: Identify and involve relevant stakeholders			
	GP2.8: Monitor and control the process			
	GP2.9: Objectively evaluate adherence			
	GP2.10: Review status with higher management			
CL 3 (Defined)	GP3.1: Establish a defined process			
	GP3.2: Collect Improvement Information			
CL 4 (Quantitatively managed)	GP4.1: Establish quantitative objectives for the process			
	GP4.2: Stabilize subprocess performance			
CL 5	GP5.1: Ensure continuous process improvement			
(Optimizing)	GP5.2: Correct root causes of problems			

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4

Improving a Process Area

	SP x.y-1 (GP1.1)	SP x.y-2 & GP1.1 ~ GP2.10	SP x.y-3 & GP3.1 & GP3.2	GP4.1 & GP4.2	GP5.1 & GP5.2
CL 1	S				
CL 2	S	S			
CL 3	S	S	S		
CL 4	S	S	S	S	
CL 5	S	S	S	S	S

Advanced practices, SP x.y-2 and SP x.y-3, exist in only the Engineering PAs.

: Base practices: SP x.y - 1



Staged Representation

25 process areas

Level	Facus	Process Areas
5 Optimizing	Continuous process improvement	Organizational Innovation and Deployment (ORGS) Causal Analysis and Resolution (SLPE)
4 Quantitatively Managed	Quantitative management	Organizational Process Performance (08/64) Countriative Project Management (PRJR)
3 Delined	Process standardization (IPPD) (83) (IPPD)	10. Rink Management (PRJS) 11. Integrated Teaming (PRJS) 12. Integrated Supplier Management (PRJ7) 13. Decision Bratysis and Revolution (SUP4)
2 Nanaged	Basic project management	1. Requirements Management (SNO1) 2. Project Parenting (PRJ*) 3. Project Menitoring and Control (PRJ2) 4. Supplier Agreement Management (PRJ2) 5. Measurement and Analysis (SUP3) 6. Process and Product Quality Assurance (SUP3) 7. Configuration Management (SIP9) 9.
1 Initial		

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Example: RD (Specific Practices)

Continuous representation SG 1 Develop Customer Requirements SP 1.1-1 Collect Stakeholder Needs SP 1.2-1 Develop the Customer Requirements SG 2 Develop Product Requirements SP 2.1-1 Establish Product & Product-Component Requirements SP 2.2-1 Allocate Product-Component Requirements SP 2.3-1 Identify Interface Requirements SP 3.1-1 Establish Operational Concepts and Scenarios SP 3.2-1 Establish a Definition of Required Functionality SP 3.3-1 Analyze Requirements SP 3.4-3 Analyze Requirements to Achieve Balance

SP 3.5-2 Validate Requirements with Comprehensive

Staged representation

SG 1 Develop Customer Requirements

SP 1.1 Elicit Needs

SP 1.2 Develop the Customer Requirements

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SP 3.5 Validate Requirements with Comprehensive Methods

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SP 3.5-1 Validate Requirements

12



Example: RD (Generic Practices)

Continuous representation

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

GG 4 Institutionalize a Quantitatively Managed Process

- GP 4.1 Establish Quantitative Objectives for the Process
- GP 4.2 Stabilize Subprocess Performance

GG 5 Institutionalize an Optimizing Process

- GP 5.1 Ensure Continuous Process Improvement
- GP 5.2 Correct Root Causes of Problems

Staged representation

GG2 Institutionalize a Managed Process

GP 2.1 (CO 1) Establish an Organizational Policy

GP 2.2 (AB 2) Plan the Process

GP 2.3 (AB 3) Provide Resources

GP 2.4 (AB 4) Assign Responsibility

GP 2.5 (AB 5) Train People

GP 2.6 (AB 5) Manage Configurations

GP 2.7 (DI 1) Identify and Involve Relevant Stakeholders

GP 2.8 (DI 2) Monitor and Control the Process

GP 2.9 (VE 1) Objectively Evaluate Adherence

GP 2.10 (VE 2) Review Status with Higher Level Management

GG 3 Institutionalize a Defined Process

GP 3.1 (AB 1) Establish a Defined Process

GP 3.2 (DI 3) Collect Improvement Information

13

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Equivalent Staging (Continuous → Staged)

	Staged representations Process Area		G.1 CL2	CL3	CL4	CER
ML2	Project Planning Project Planning and Comod Supplier Agrormed Management Requirement Management Configuration Management Configuration Management Process and Product Quality Resource Mountement and Analysis	PF- PHIC SAMA RESUM PF-SA CM MA	Target Profile 2			
ML3	Cognitional Process Terror Cognitional Process Terror Cognitional Process Terror Cognitional Surving Cognitional Cognitional Surving Cognition Co	OPP OPE OF	Target Pysille 3		One or more than see PAs	One or reason than one Phil
MLA	Organizational Process Performance Occanization Propertition agreement	DIM.	Turget Per	me e		
MLS	Cognitational Innovation and Deployment Council Analysis and Emolution	CAR	TorquitPro	ene s		

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Common Terminology with Special Meaning (26)

- Establish and maintain (in goals and practices)
 - formulated, documented, and used throughout the organization

Customer

- is the party (individual, project, or organization) responsible for <u>accepting the</u> <u>product</u> or <u>authorizing payment</u>.
- external to the project, but not necessarily external to the organization.
- subset of stakeholders

Stakeholder

- a group or individual that is affected by or in some way accountable for the outcome of an undertaking.
- example: project managers, suppliers, customers, end users, and others.
- o Relevant stakeholder
 - designate a stakeholder that is <u>identified</u> for involvement in specified activities and is <u>included</u> in an appropriate plan.

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15



Product

- Any tangible output or service that is a result of a process and that is intended for deliver to a customer or end user.
- Product is a work product that is delivered to the customer.

Work product

- Any artefacts produced by process. e.g., files, documents, parts of product, services, processes, specifications, and invoices.
- Manufacturing process, training process, and disposal process for the product. <u>A work product need not be engineered or part of the end</u> product.

Product component

- Lower level components of the product.
- Any work product that must be engineered.



- Quality and process-performance (QPP) objectives
 - Covers <u>objectives and requirements</u> for product quality, service quality, and process performance.
 - Process measurements (e.g., effort, cycle time, and defect removal effectiveness) and
 - Product measurements (e.g., reliability and defect density).

o Organization's set of standard processes (OSSPs)

- Contains the definitions of the processes (process descriptions) that guide all activities in an organization.
- These process description covers the fundamental process elements (their relationships to each other) that must be incorporated into the defined process. (process architecture)



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17



CMMI-Specific terminology (14)

- Organization's process asset library (OPAL)
 - <u>Examples of process-related documentation:</u> policies, defined processes, checklists, lessons-learned documents, templates, standards, procedures, plans, and other artefacts.
- Organization's measurement repository (OMR)
 - Examples of process and work product data: estimated size of work products, actual effort expended, and actual costs; peer review efficiency and coverage statistics; and the number of and severity of defects.
- Organization process assets (OPAs)
 - Org's process assets at CMMI; (5+2)
 - Organization's set of standard processes, including the <u>process architectures</u> and <u>process elements</u>.
 - Descriptions of life-cycle models approved for use
 - Guidelines and criteria for tailoring the organization's set of standard processes
 - Organization's measurement repository
 - Organization's process asset library
 - Organizational process performance baselines
 - Organizational process performance models



4. Capability Levels and Generic Model Components



Introduction to CMMI

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19



Capability Level 0: Incomplete

- o Process is either <u>not performed</u> or <u>partially performed</u>.
- o One or more of the specific goals of the PA are not satisfied.

A process performance is not step function (by Ho-Won).



Capability Level 1: Performed

- All specific goals of the PA are satisfied.
- Produce identified output work products using identified input work products.
- o The process may be unstable and inconsistently implemented.
- o **GG 1:** Achieve specific goals.
- o **GP 1:** Perform <u>base practices</u> of the PA to develop work products and provide services to achieve the specific goals of the PA.



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21



Capability level 2: Managed

- o A managed process is a performed process that
 - is also planned and executed in accordance with policy;
 - employs skilled people having adequate resources;
 - involves relevant stakeholders:
 - is monitored, controlled, and reviewed;
 - is evaluated for adherence to its process description.
- Processes are managed to achieve <u>process objectives</u> such as cost, schedule, and quality.

Process objectives: cost, schedule, and quality.



GG 2: Institutionalize a Managed Process

o GP 2.1 Establish an Organization Policy

- Establish and maintain an organization policy for planning and performing the process.
- The purpose is to define organizational <u>expectations for the process</u> and make these expectations visible to those affected in the organization.

GP 2.2 Plan the Process

- Establish and maintain the plan for performing the process.
- The purpose is:
 - to determine what is needed to perform the process and achieve the established objective;
 - to prepare a plan for performing the process;
 - to prepare a process description;
 - to get agreement on the plan from relevant stakeholders.
- Plan may be from other processes (PMC plan PP

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23



o GP 2.3 Provide Resources

- Provide adequate resources for performing the process, developing the work products, and providing the services of the process.
 - Resources: Funding, people, tools, facilities

o GP 2.4 Assign Responsibility

 Assign responsibility and authority for performing the process, developing the work product, and providing the services of the process.

o GP 2.5 Train People

- Train the people performing or supporting the process as needed.
- Refer to the Organizational Training (OT)

GP 2.6 Manage Configuration

 Place designated work products of the process (identified in the plan for performing the process) under appropriate levels of CM.



GP 2.7 Identify and Involve Relevant Stakeholders

 The purpose is to establish and maintain the expected involvement of stakeholders during the execution of the process, and to ensure that interactions necessary to the process are accomplished.

o GP 2.8 Monitor and Control the Process

 Monitor and control the process <u>against the plan</u> for performing the process and <u>take appropriate corrective action</u>.

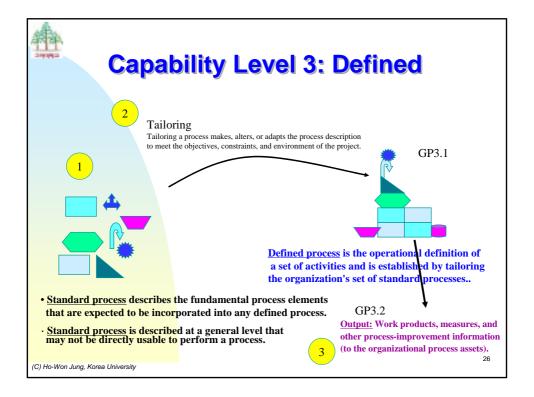
o GP 2.9 Objectively Evaluate Adherence

 Objectively evaluate adherence of the process against its process description, standard, and procedures, and address non-compliances.

o GP 2.10 Review Status with Higher Level Management

- Review the activities, status, and results of the process with higherlevel management and <u>resolve issues</u>.
- The purpose is to provide higher-level management with the appropriate visibility into the process.

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GG 3: Institutionalize a Defined Process

- GP 3.1 Establish a Defined Process.
 - Establish and maintain the description of a defined process that is tailored from the OSSPs (Organization's Set of Standard Processes).
 - · Variability of performing process across the organization is reduced
 - Process asset, data, and learning can be shared effectively.
- GP 3.2 Collect Improvement Information.
 - Collect work products, measures, measurement results, and improvement
 information derived from planning and performing the process to support the
 future use and improvement of the OPPAs (Organization's Processes and
 Process Assets).
 - The information and artefacts are stored in the organization's measurement repository and the organization's process library.
 - The process and product measures are primarily those that are defined in the common set of measures for the OPPAs.

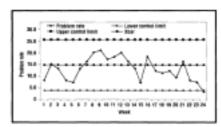
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27



Capability Level 4: Quantitatively Managed

- Quantitatively managed process is a defined process that
 - is controlled using <u>statistical and</u> <u>other quantitative techniques.</u>



- Predictability (stability): variation within control limits.
- · Capable process: meet specification, objective, or requirement

Do not confuse the differences between "predictability" and "prediction".

OPP: Quality and process-performance



Source of process performance variation

[Total variation] = [Special cause variation]



[Common cause variation]

Special cause = Assignable cause

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29



Special (assignable) Causes

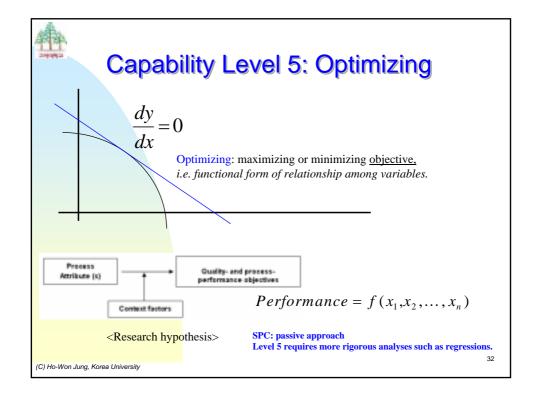
- Preventable variations
- Events from not part of the normal process.
- Removing all special causes results in a <u>stable and predictable process</u>.
- Examples of special causes of variation
 - Failures to follow the process (lack of process compliance)
 - Changes in the quality of raw materials (uncontrolled inputs to the subprocess)
 - · Changes to work environment
 - Inadequately trained people
 - Schedule pressure
 - Tool failures
 - Changes in methods used
 - Inappropriate sampling or grouping of data



GG 4: Institutionalize a Quantitatively Managed Process

- o GP 4.1 Establish Quantitative Objectives for the Process.
 - Establish and maintain <u>quantitative objectives for the process</u>.
 - The purpose is to determine and obtain <u>agreement</u> from <u>relevant</u> <u>stakeholders</u> about specific quantitative objectives for the process.
- o GP 4.2 Stabilize Subprocess Performance.
 - Stabilize the performance of one or more subprocesses to determine the ability of the process to achieve the established QPP objectives.
 - Refer to the QPM SG 2 (Statistically Manage Subprocess Performance)

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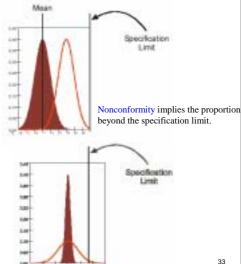


Continuous Process Improvement

If the process is both stable and capable, then the organization should continuously seek ways to improve the variation (narrower control limits) and improve the capability of the process

 Changing the mean results in reducing the proportion of the nonconformities.

 Reducing the variation results in reducing the proportion of the nonconformities.



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- o GP 5.1 Ensure Continuous Process Improvement.
 - Ensure continuous improvement of the process in fulfilling the relevant business objectives of the organization.
 - The purpose is to select and systematically deploy process and technology <u>improvements</u> that contribute to <u>meeting</u> "<u>established</u> <u>quality and PI objectives</u>".
- o GP 5.2 Correct Root Causes of Problems.
 - Identify and correct the root causes of defects and other problems in the process.
 - The purpose is to analyze defects and other problems, to correct the root causes of these types of defects and problems, and to prevent these defects and problems from occurring in the future.