Failure Mode and Effects Analysis (FMEA)

UNDERSTANDING THE FUNDAMENTAL DEFINITIONS AND CONCEPTS OF FMEAS

Definition of FMEA

Failure Mode and Effects Analysis (FMEA) is a method designed to:

- Identify and fully understand potential failure modes and their causes, and the effects of failure on the system or end users, for a given product or process.
- Assess the risk associated with the identified failure modes, effects and causes, and prioritize issues for corrective action.
- Identify and carry out corrective actions to address the most serious concerns.

What is FMEA?

An FMEA is an engineering analysis

- done by a cross-functional team of subject matter experts
- that thoroughly analyzes product designs or manufacturing processes
- early in the product development process.
- Finds and corrects weaknesses before the product gets into the hands of the customer.

What is FMEA?

An FMEA should be the guide to the development of a complete set of actions that will reduce risk associated with the system, subsystem, and component or manufacturing/assembly process to an acceptable level.

What is FMEA?

- Performing an FMEA just to fill a checkbox in the Product Development Process and then filing it away, never to be seen again, is a waste of time and adds no value.
- If not for use as guidance through the development process, why waste the time and resources to do it in the first place?
- If effectively used throughout the product life cycle, it will result in significant improvements to reliability, safety, quality, delivery, and cost.

Primary Objective of FMEA

The primary objective of an FMEA is to improve the design.

- For System FMEAs, the objective is to improve the design of the system.
- For Design FMEAs, the objective is to improve the design of the subsystem or component.
- For Process FMEAs, the objective is to improve the design of the manufacturing process.

Primary Objective of FMEA

There are many other objectives for doing FMEAs, such as:

- identify and prevent safety hazards
- minimize loss of product performance or performance degradation
- improve test and verification plans (in the case of System or Design FMEAs)
- improve Process Control Plans (in the case of Process FMEAs)
- consider changes to the product design or manufacturing process
- identify significant product or process characteristics
- develop Preventive Maintenance plans for in-service machinery and equipment
- develop online diagnostic techniques

Types of FMEAs

The three most common types of FMEAs are:

- System FMEA
- Design FMEA
- Process FMEA

System FMEA

Analysis is at highest-level analysis of an <u>entire</u> <u>system</u>, made up of various subsystems.

The **focus** is on system-related deficiencies, including

- system safety and system integration
- interfaces between subsystems or with other systems
- interactions between subsystems or with the surrounding environment
- single-point failures (where a single component failure can result in complete failure of the entire system)

System FMEA

The **focus** (continued)

- functions and relationships that are *unique* to the system as a whole (i.e., do not exist at lower levels) and could cause the overall system not to work as intended
- human interactions
- service

Some practitioners separate out human interaction and service into their own respective FMEAs.

Design FMEA

Analysis is at the subsystem level (made up of various components) or component level.

The **Focus** is on product design-related deficiencies, with emphasis on

- improving the design
- ensuring product operation is safe and reliable during the useful life of the equipment.
- interfaces between adjacent components.

Design FMEA usually assumes the product will be manufactured according to specifications.

Process FMEA

Analysis is at the manufacturing/assembly process level.

The **Focus** is on manufacturing related deficiencies, with emphasis on

- Improving the manufacturing process
- ensuring the product is built to design requirements in a safe manner, with minimal downtime, scrap and rework.
- manufacturing and assembly operations, shipping, incoming parts, transporting of materials, storage, conveyors, tool maintenance, and labeling.

Process FMEAs most often assume the design is sound

FMEA Definitions and Examples

Generic FMEA Worksheet

		Potential	Potential	λ.	Potential	nce	Current	Current	uc			Responsible Person	Actions Taken	l		ised kings	
Item	Function	Failure Mode	Effect(s) of Failure	Severit	Potential Cause(s) of Failure	Occurrence	Design Controls (Prevention)	Design Controls (Detection)	Detection	RPN	Recommended Action(s)	Target Completion Date	Effective Completion Date	Severity	Occurrence	Detection	RPN
1	2	3	4	5	6	7	8	8	9	10	11)		12)				

Item 1

An "item" is the focus of the FMEA project.

- For a System FMEA this is the system itself.
- For a Design FMEA, this is the subsystem or component under analysis.
- For a Process FMEA, this is usually one of the specific steps of the manufacturing or assembly process under analysis, as represented by an operation description.

Example: Bicycle hand brake subsystem

All-Terrain System Hierarchy

(with components for Hand Brake Subsystem)

- 1.0 All-Terrain Bicycle System
 - 1.1 Frame Subsystem
 - 1.2 Front Wheel Subsystem
 - 1.3 Rear Wheel Subsystem
 - 1.4 Sprocket-Pedal Subsystem
 - 1.5 Chain-Derailleur Subsystem
 - 1.6 Seat Subsystem
 - 1.7 Handle Bar Subsystem
 - 1.8 (Hand Brake Subsystem
 - 1.8.1 Brake Lever
 - 1.8.2 (Brake Cable
 - 1.8.3 Brake Pads
 - 1.8.4 Brake Calliper
 - 1.9 Suspension Subsystem

Item identification for All-Terrain System FMEA

> Item identification for All-Terrain Hand Brake Design FMEA

> > Item identification for All-Terrain Cable Design FMEA

Function (2)

A "function" is what the item or process is intended to do, usually to a given standard of performance or requirement.

- For Design FMEAs, this is the primary purpose or design intent of the item.
- For Process FMEAs, this is the primary purpose of the manufacturing or assembly operation.
- Functions are typically described in a verb-noun format.
- There can be many functions for each item or operation.

Example: Provides the correct level of friction between brake pad assembly and wheel rim to safely stop bicycle in the required distance, under all operating conditions.

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Failure Mode 3

The term "failure mode" combines two words that both have unique meanings.

- The Concise Oxford English Dictionary defines the word "failure" as the act of ceasing to function or the state of not functioning.
- "Mode" is defined as a way in which something occurs

Failure Mode 3

- A "failure mode" is the manner in which the item or operation potentially fails to meet or deliver the intended function and associated requirements.
- may include failure to perform a function within defined limits
- inadequate or poor performance of the function
- intermittent performance of a function
- and/or performing an unintended or undesired function Example: Insufficient friction delivered by hand brake subsystem between brake pads and wheels during heavy rain conditions.

Item/Function	Potential Failure Mode	Potential Effect(s) of Failure
Hand Brake S/S:	Insufficient	Bicycle wheel
Provides the	friction delivered	does not slow
correct level of	by hand brake	down when the
friction between	subsystem	brake lever is
brake pad	•	pulled
assembly and	between brake	potentially
wheel rim to	pads and wheels	resulting in
safely stop bicycle	during heavy rain	accident.
in the required	conditions.	
distance, under all		
operating		
conditions.		

TRUNCATED

Effect 4

An "effect" is the consequence of the failure on the system or end user.

- This can be a single description of the effect on the toplevel system and/or end user, or three levels of effects (local, next-higher level, and end effect)
- For Process FMEAs, consider the effect at the manuf. or assembly level, as well as at the system or end user.
- There can be more than one effect for each failure mode. However, typically the FMEA team will use the most serious of the end effects for the analysis.

Example: Bicycle wheel does not slow down when the brake lever is pulled potentially resulting in accident.

Item/Function	Potential Failure Mode	Potential Effect(s) of Failure	S E V
Hand Brake S/S:	Insufficient	Bicycle wheel does	10
Provides the	friction delivered	not slow down	
correct level of	by hand brake	when the brake	
friction between	subsystem	lever is pulled	
brake pad assembly	between brake		
and wheel rim to	pads and wheels	potentially	
safely stop bicycle	during heavy	resulting in	
in the required	rain conditions.	accident.	
distance, under all			
operating			
conditions			

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Severity 5

- "Severity" is a ranking number associated with the most serious effect for a given failure mode
- based on the criteria from a severity scale.
- a relative ranking within the scope of the specific FMEA
- determined without regard to the likelihood of occurrence or detection.

Example: 10

	Item/Function	Potenti Failur Mode	e	Potential Effect(s) of Failure	S E	Potential Cause(s) of Failure	0 c c	
	Bicycle System: The bicycle must lation Criteria on Product	Does no stop in	d e	Potential accident or injury to bicycle operator	10	nsufficient friction delivered by hand brake subsystem between brake	5	
	ration and/or involves ithout warning.	Rank		without warning.		pads and wheels during heavy rain conditions.		
operation and/or involves n with warning.		9				Brake system mis- adjusted by bicycle	3	NCATED
e, d	oes not affect safe	8				user		NCA

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Suggested DFMEA Severity Evaluation Criteria

Effect	Criteria: Severity of Effect on Product (Customer Effect)	Rank					
Failure to Meet Safety and/or	Potential failure mode affects safe vehicle operation and/or involves noncompliance with government regulation without warning.	10					
Regulatory Requirements	Potential failure mode affects safe vehicle operation and/or involves noncompliance with government regulation with warning.	9					
Loss or Degradation of	Loss of primary function (vehicle inoperable, does not affect safe vehicle operation).						
Primary Function	Degradation of primary function (vehicle operable, but at reduced level of performance).						
Loss or Degradation of	Loss of secondary function (vehicle operable, but comfort / convenience functions inoperable).						
Secondary Function	Degradation of secondary function (vehicle operable, but comfort / convenience functions at reduced level of performance).	5					
	Appearance or Audible Noise, vehicle operable, item does not conform. Defect noticed by most customers (> 75%).						
Annoyance	Appearance or Audible Noise, vehicle operable, item does not conform. Defect noticed by many customers (50%).						
	Appearance or Audible Noise, vehicle operable, item does not conform. Defect noticed by discriminating customers (< 25%).						
No Effect	No discernible effect.	1					

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Cause 6

A "cause" is the specific reason for the failure, preferably found by asking "why" until the root cause is determined.

- For Design FMEAs, the cause is the *design deficiency* that results in the failure mode.
- For Process FMEAs, the cause is the *manufacturing or assembly deficiency* that results in the failure mode.
- at the component level, cause should be taken to the level of failure mechanism.
- if a cause occurs, the corresponding failure mode occurs.
- There can be many causes for each failure mode.

Example: Cable breaks

Item/Function	Potential Failure Mode	Potential Effect(s) of Failure	S E V	Potential Cause(s) of Failure	0 C C
Hand Brake S/S: Provides the correct level of friction between brake pad assembly and wheel	by hand brake subsystem	Bicycle wheel does not slow down when the brake lever is pulled	10	Cable Binds due to inadequate lubrication or poor routing	4
rim to safely stop bicycle in the required distance, under all operating	pads and wheels during heavy rain conditions.	potentially resulting in accident.		External foreign material reduces friction	2
conditions.				Cable breaks	6

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Occurrence (7)

"Occurrence" is a ranking number associated with the likelihood that the failure mode and its associated cause will be present in the item being analyzed.

- For System and Design FMEAs, consider the likelihood of occurrence during the design life of the product.
- For Process FMEAs consider the likelihood of occurrence during production.
- based on the criteria from the corresponding occurrence scale.
- has a relative meaning rather than absolute value, determined without regard to the severity or likelihood of detection.

Example: 6

		Item/Function	Potential Failure Mode	Potential Effect(s) of Failure	S E V	Potential Cause(s) of Failure	0 C	
Sugge	ested DFMEA Occur	-	-	Potential accident or injury to bicycle or	10	Insufficient friction delivered by hand brake subsystem between brake pads and wheels	5)	
Likelihood of Failure	Criteria: Occui (Design Life/Reliabi	rrence of Cause ility of Item/Vehicle	e) Ra	nk ^{g.}		during heavy rain conditions.		
Very High	New technology/new design with no history.			0		Brake system mis-	3 0	
	Failure is inevitable with ne or change in duty cycle/ope		n, g			adjusted by bicycle user	TRUNCATED	
High	Failure is likely with new de change in duty cycle/operat		8	NCAT	ED		T X	
	Failure is uncertain with new change in duty cycle/operate		, or 7					
	Frequent failures associated design simulation and testir	_	1 6					
Moderate	Occasional failures associate design simulation and testin	_	in s	y		•		
	Isolated failures associated design simulation and testir	•	4					
Low	Only isolated failures associ design or in design simulation		' (3	y				
Low	No observed failures associa design or in design simulation		2	Tr		ustration is from the boo		
Very Low	Failure is eliminated throug	h preventative control.	1		ırı S.	Carlson, published by Jo	onn wney	& Sons, © 2012

Controls (8)

"Controls" are the methods or actions <u>currently</u> planned, or are already in place, to reduce or eliminate the risk associated with each potential cause.

- Controls can be the methods to prevent or detect the cause during product development, or actions to detect a problem during service before it becomes catastrophic.
- There can be many controls for each cause.

Prevention-type Controls

- For System or Design FMEAs, prevention-type design controls describe how a cause, failure mode, or effect in the product design is *prevented* based on current or planned actions
- they are intended to reduce the likelihood that the problem will occur, and are used as input to the occurrence ranking.

Example: Cable material selection based on ANSI #ABC.

Detection-type Controls

- For System or Design FMEAs, detection-type designs controls describe how a failure mode or cause in the product design is *detected*, based on current or planned actions before the product design is released to production, and are used as input to the detection ranking.
- They are intended to increase the likelihood that the problem will be detected before it reaches the end user.

Example: Bicycle system durability test # 789

Item/Function	Potential Failure Mode	Potential Effect(s) of Failure	S E V	Potential Cause(s) of Failure	0 C C	Current Design Controls (Prevention)	Current Design Controls (Detection)	D E T	
friction between	friction delivered by hand brake subsystem	down when the brake lever is		Cable Binds due to inadequate lubrication or poor routing	4	Hand Brake Design Guide #123	Bicycle system durability test # 789	2	
brake pad assembly and wheel rim to safely stop bicycle	between brake pads and wheels during heavy rain conditions.	pulled potentially resulting in accident.		External foreign material reduces friction	2			3	
in the required distance, under all operating conditions.				Cable breaks	6	Cable material selection based on ANSI #ABC.	Bicycle system durability test # 789	4	

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Detection (9)

- "Detection" is a ranking number associated with the best control from the list of detection-type controls, based on the criteria from the detection scale.
- considers the likelihood of detection of the failure mode/cause, according to defined criteria.
- a relative ranking within the scope of the specific FMEA
- determined without regard to the severity or likelihood of occurrence.

Example: 4

-								
	Item/Function	Potential Failure Mode	Potential Effect(s) of Failure	S E V	Potential Cause(s) of Failure	0 C C	Current Design Contr (Prevention)	ols Current Design Controls (Detection)
] F r t i	Bicycle System: The bicycle must provide safe and reliable transportation, including safe DISCOURGE OF THE PROPERTY OF THE PR	distance	Potential accident or injury to bicycle operator without etection		Insufficient friction delivered by hand brake subsystem between brake pads and wheels during heavy rain raluation Cri		All-Terrain braking system design guid (document # 123)	' 1 ' 11
Opportunity for Detection	Likelihood		teria: tion by De	esig	n Control	Rank	Likelihood of Detection	1. Bicycle system 9 durability teat # 789
No detection opportunity	No current design control; Cannot detect or is not analyzed.				lyzed.	10	Absolute Uncertainty	2 dicycle system performance testing to design
Not likely to detect at any stage	Design analysis/detection controls have a weak detection capability; Virtual Analysis (e.g. CAE, FEA, etc.) is <u>not correlated</u> to expected actual operating conditions.					9	Very Remote	requirements
	Product verification/validation after design freeze and prior to launch with <u>pass/fail</u> testing (Sub-system or system testing with acceptance criteria e.g. ride & handling, shipping evaluation, etc.)						Remote	
Post Design Freeze and prior to launch	Product verification/validation after design freeze and prior to launch with <u>test to failure</u> testing (Sub-system or system testing until failure occurs, testing of system interactions, etc.)						Very l w	
	Product verification/validation after design freeze and prior to launch with <u>degradation</u> testing (Sub-system or system testing after durability test e.g. function check).						Low	
_	Product validation (re prior to design freeze performance, functio	using pass/fa	il testing (e.g.			5	Moderately	
Prior to Design	Product validation (re	liability testin	g, developme	nt or	validation tests)			

Moderately High

High

4

3

Freeze

etc.).

before/after values, etc.)

prior to design freeze using test to failure (e.g. until leaks, yields, cracks,

Product validation (reliability testing, development or validation tests)

prior to design freeze using degradation testing (e.g. data trends,

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Risk Priority Number (RPN)(10)

"RPN" is a numerical ranking of the risk of each potential failure mode/cause, made up of the arithmetic product of the three elements:

- severity of the effect
- likelihood of occurrence of the cause
- likelihood of detection of the cause.

Example: 240 (10 x 6 x 4)

Limitations of RPN

RPN is not a perfect representation of the risk associated with a failure mode and associated cause.

- subjective
- not continuous

High severity must be considered regardless of RPN value

Recommended Actions (11)



"Recommended actions" are the tasks recommended by the FMEA team to reduce or eliminate the risk associated with potential causes of failure. They should consider

- existing controls
- relative importance (prioritization) of the issue
- cost and effectiveness of the corrective action.
- there can be many recommended actions for each cause.

Example: Require cable DFMEA/PFMEA from cable supplier approved by All-Terrain FMEA team.

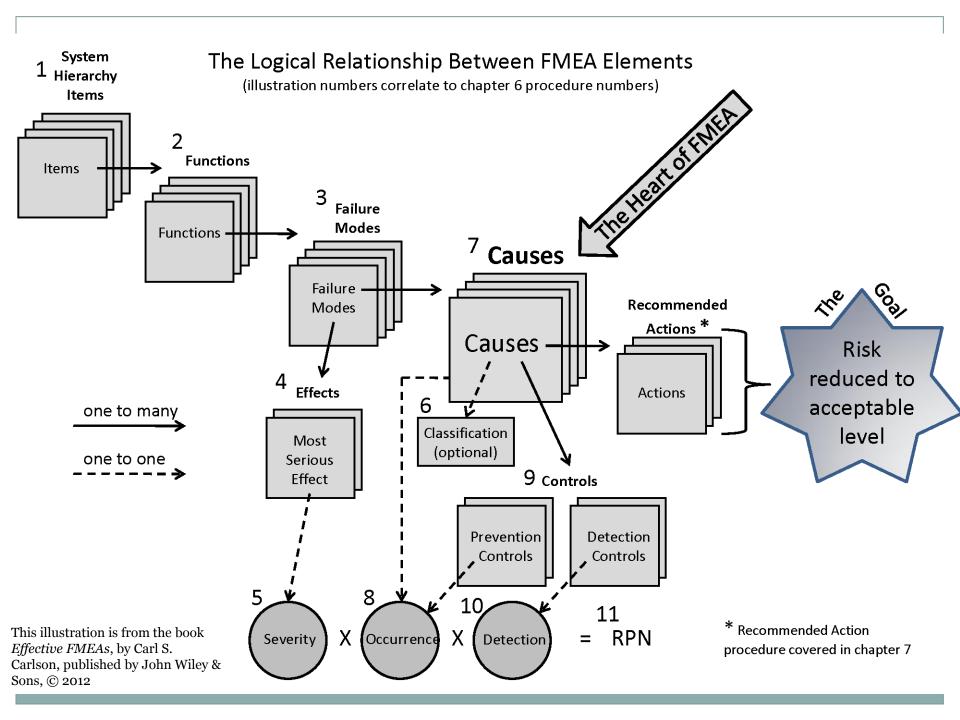
Item/Function	Potential Failure Mode	Potential Effect(s) of Failure	S E V	Potential Cause(s) of Failure	0 C C	Current Design Controls (Prevention)	Current Design Controls (Detection)	D E T	R P N	Recommended Actions
	friction delivered by hand brake	Bicycle wheel does not slow down when the brake lever is	10	Cable Binds due to inadequate lubrication or poor routing		Hand Brake Design Guide #123	Bicycle system durability test # 789	2	80	Redesign hand brake cable routing to reduce friction and make system insensitive to lubrication degradation
assembly and wheel rim to	pads and wheels during heavy	pulled potentially resulting in accident.								Modify bicycle durability testing to include periodic brake cable checks for binding
bicycle in the required distance, under				External foreign material reduces friction	2			3	60	
all operating conditions.				Cable breaks		Cable material selection based on ANSI #ABC.	Bicycle system durability test # 789	4	240	Require cable DFMEA/PFMEA from cable supplier approved by All-Terrain FMEA team.
										Based on results of Cable DFMEA, develop cable strength test and modify cable design to improve strength
				Brake Lever breaks		Hand Brake Design Guide #123	Bicycle system durability test # 789	1	10	
				Selected brake pad material does not apply required friction to wheel	2			2	40	

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Actions Taken (12)

- "Action Taken" is the specific action that is implemented to reduce risk to an acceptable level.
- it should correlate to the specific recommended action
- and is assessed as to effectiveness by a revised severity, occurrence, detection ranking, and corresponding revised RPN.

Example: Cable supplier completed DFMEA/PFMEA and approved by All-Terrain team



Is that all there is to FMEA?

- If FMEA were only an exercise in "filling out a form" then the definitions would be all you need to know.
- There is much more to learn about FMEAs!

What else is needed?

- FMEA has the *potential* to anticipate and prevent problems, reduce costs, shorten product development times, and achieve safe and highly reliable products and processes.
- To obtain the best possible results from FMEA, companies need to focus on key success factors.

FMEA Success Factors

- understanding the fundamentals and procedure of FMEAs, including the concepts and definitions
- preparation steps for each FMEA project
- applying lessons learned and quality objectives
- providing excellent facilitation
- and implementing an effective company-wide FMEA process.

Implementing FMEA success factors will uniformly ensure FMEAs achieve safe, reliable and economical products and processes.

Preparation Steps for Each FMEA Project

- Determine the scope of the FMEA project
- Make the scope visible and get consensus on boundaries
- Assemble the right FMEA team (not done by one or two people)
- Establish ground rules and assumptions
- Gather information
- Prepare for the FMEA meetings

Applying Lessons Learned & Quality Objectives

Much is learned by observing the mistakes companies have made in doing FMEAs. Based on the experience of over two thousand FMEAs and working with hundreds of companies in a wide variety of applications, certain common mistakes show up repeatedly.

- What are the primary ways that FMEAs can be done wrongly (mistakes made)
- What are the leading factors that make for effective FMEAs (quality objectives)?

Providing excellent facilitation

- FMEA facilitation is a different subject than FMEA methodology.
- To be successful, FMEA leaders need to develop expert facilitation skills
 - Brainstorming
 - Encouraging Participation
 - O Active Listening
 - Controlling Discussion
 - O Making Decisions
 - Conflict Management
 - Managing Level of Detail
 - O Managing Time
 - Unleashing Team Creativity

Implementing an effective FMEA process

A *company-wide FMEA process* is the entire set of systems and tasks essential to support development of high-reliability products and processes through timely accomplishment of well-done FMEAs.

- Management support for strategy and resources
- Roles and responsibilities
- Management review of high risk issues
- FMEA quality audits
- Execution of FMEA recommended actions
- Feedback loop to incorporate lessons learned

In Summary...

- Everyone wants to support the accomplishment of safe and trouble-free products and processes while generating happy and loyal customers.
- When done correctly, FMEA can anticipate and prevent problems, reduce costs, shorten product development times, and achieve safe and highly reliable products and processes.

FMEA PROBLEMS

AND SOLUTIONS

Which of the following are true statements about FMEA? (Select all that apply)

- 1. An FMEA is an engineering analysis done by the most knowledgeable person on the engineering team.
- 2. Part of the FMEA is to identify and carry out corrective actions to address the most serious concerns.
- 3. The primary objective of an FMEA is to understand the design.
- 4. Risk assessment is not part of the FMEA procedure.

Which of the following are true statements about FMEA? (Select all that apply.)

- 1. An FMEA is an engineering analysis done by the most knowledgeable person on the engineering team. (False. An FMEA is an engineering analysis done by a cross-functional team of subject-matter experts.)
- 2. Part of the FMEA is to identify and carry out corrective actions to address the most serious concerns. (True)
- 3. The primary objective of an FMEA is to understand the design. (False. The primary objective of an FMEA is to *improve* the design.)
- 4. Risk assessment is not part of the FMEA procedure. (False. Risk assessment is an integral part of the FMEA procedure.)

Indicate whether each statement about the application of FMEA is true or false.

- 1. One of the uses of FMEA is to improve the reliability of the product.
- 2. One of the uses of FMEA is to improve the safety of the product.
- 3. FMEAs can be used to improve the quality of the manufacturing process.
- 4. One of the primary applications of FMEA is to fix field problems.

Indicate whether each statement about the application of FMEA is true or false.

- 1. One of the uses of FMEA is to improve the reliability of the product. (True)
- 2. One of the uses of FMEA is to improve the safety of the product. (True)
- 3. FMEAs can be used to improve the quality of the manufacturing process. (True)
- 4. One of the primary applications of FMEA is to fix field problems. (False)

In an FMEA, which of the following is true about a "function"? (Select all that apply)

- 1. A "function" is what the item is intended to do, without respect to any standard of performance.
- 2. A "function" is what the item is intended to do, usually to a given standard of performance.
- 3. There is one function for each item in an FMEA.
- 4. The function description in an FMEA must include the consequence or impact on the end user.

In an FMEA, which of the following is true about a "function"? (Select all that apply)

- 1. A "function" is what the item is intended to do, without respect to any standard of performance. (False. A function description needs to include the standard of performance.)
- 2. A "function" is what the item is intended to do, usually to a given standard of performance. (True)
- 3. There is one function for each item in an FMEA. (False. There can be many functions for an item.)
- 4. The function description in an FMEA must include the consequence or impact on the end user. (False. An effect must include the consequence or impact on the end user, not a function.)

In an FMEA, which of the following is true about a "failure mode"? (Select all that apply)

- 1. A "failure mode" is the specific reason for the failure.
- 2. A "failure mode" is the manner in which the item or assembly could fail to meet the intended function and its requirements
- 3. In an FMEA, there is one failure mode for each function.
- 4. The failure mode description in an FMEA must include the consequence or impact on the end user.

In an FMEA, which of the following is true about a "failure mode"? (Select all that apply)

- 1. A "failure mode" is the specific reason for the failure. (False. A "failure mode" is the manner in which the item or assembly could fail to meet the intended function and its requirements.)
- 2. A "failure mode" is the manner in which the item or assembly could fail to meet the intended function and its requirements. (True)
- 3. In an FMEA, there is one failure mode for each function. (False. There can be many failure modes for each function.)
- 4. The failure mode description in an FMEA must include the consequence or impact on the end user. (False. An effect must include the consequence or impact on the end user, not a failure mode.)

In an FMEA, which of the following is true about a "control"? (Select all that apply)

- 1. A "control" is the specific recommendation by the FMEA team to control the risk associated with the cause of failure.
- 2. A "control" needs to be taken to the level of root cause of the failure.
- 3. There are often two types of controls identified in an FMEA: prevention-type controls and detection-type controls.
- 4. "Controls" are the methods or actions that are not currently planned, but need to be done to reduce or eliminate the design-related risk associated with the cause of failure.
- 5. "Controls" are the methods or actions that are planned or currently in place to reduce or eliminate the design-related risk associated with the cause of failure.

In an FMEA, which of the following is true about a "control"? (Select all that apply)

- 1. A "control" is the specific recommendation by the FMEA team to control the risk associated with the cause of failure. (False. Controls are the methods or actions that are planned or currently in place to reduce or eliminate the design-related risk associated with the cause of failure. Recommendations need to be in the Recommended Actions column of the FMEA.)
- 2. A "control" needs to be taken to the level of root cause of the failure. (False. Causes in the FMEA need to be taken to the level of root cause, not controls.)
- 3. There are often two types of controls identified in an FMEA: prevention-type controls and detection-type controls. (True)
- 4. "Controls" are the methods or actions that are not currently planned, but need to be done to reduce or eliminate the design-related risk associated with the cause of failure. (False. Controls are methods or actions that are *planned or currently in place*.)
- 5. "Controls" are the methods or actions that are planned or currently in place to reduce or eliminate the design-related risk associated with the cause of failure. (True)

FMEA Resources

- This presentation is based on the book *Effective FMEAs*, by Carl S. Carlson, published by John Wiley & Sons, © 2012
- Information about the book and links to useful FMEA articles and aids can be found on www.effectivefmeas.com.
- If you have questions or comments about this presentation, the subject of FMEAs, or the book *Effective FMEAs*, please send an email to the author at Carl.Carlson@EffectiveFMEAs.com.

Biography

- Carl S. Carlson is a consultant and instructor in the areas of FMEA, reliability program planning and other reliability engineering disciplines, currently supporting clients of ReliaSoft Corporation.
- He has 30 years experience in reliability testing, engineering, and management positions, including manager of product reliability at General Motors.
- He co-chaired the cross-industry team that developed the commercial FMEA standard (SAE J1739, 2002 version) and was a past member of the Reliability and Maintainability Symposium (RAMS) Advisory Board.
- He holds a B.S. in Mechanical Engineering from the University of Michigan, is a senior member of ASQ and a Certified Reliability Engineer.
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