AIAG & VDA FMEA Handbook

April 4, 2019



Agenda

- Current Development Status
- Project Objective
- Leveraging the new FMEA Handbook
- Examples of major changes and their benefits
- Q&A

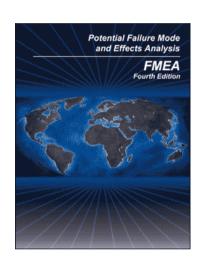


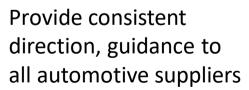
Current Development Status

- Final Draft in Approval Process
 - AIAG QSC: April 2, 2019
 - VDA QMA: May 8, 2019
 - Both approvals required to release the document
- Release of Handbook June 2019
 - Launch events in Germany and U.S.
- Availability of Training Q3 2019
 - Each Association updating training courses



AIAG VDA FMEA - Project Objective













Update to include:

- Best Practices
- Improved Examples
- Functional Safety



Importance of New Methods and Tools

- Effective FMEA risk identification never been more critical
 - Rapid growth in component/system interactions
 - Increasingly specialized technologies
 - No change in legal obligations of producers
- Effective FMEA includes:
 - Cross-functional team contributions
 - Carefully identified system boundaries
 - Thorough documentation of risks and actions



Leveraging the AIAG & VDA FMEA Handbook

- The Essential Link Between COQ and FMEA
 - You will need FMEA to make sure COQ / COPQ improvement targets are met;
 - You will need COQ / COPQ actual performance on similar products and processes to make sure the FMEA risk evaluation of the new product and/or process is realistic;
 - Not acceptable reduction in Cost of Poor Quality (COPQ) means FMEA was not effective



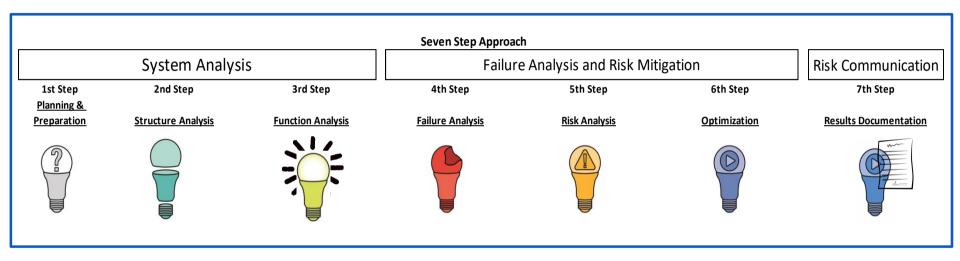
Examples of Major Changes and Benefits

- 7 Step Approach
- Supplemental FMEA MSR
- New Severity, Occurrence, Detection Tables
- PFMEA Failure Analysis
- Action Priority (AP) Tables

More Structured Approach – Leverages Lessons Learned – Prevention Driven



AIAG &VDA FMEA - 7 Step Approach



Applies to DFMEA, Supplemental FMEA – MSR, and PFMEA



Supplemental FMEA - MSR

- FMEA MSR = Monitoring and System Response
 - Supplemental approach for Design FMEA
 - Addresses Risk Analysis of "Mechatronic Systems"
 - Not previously addressed in AIAG 4th Edition FMEA
 - Describes linkages between Design FMEA and Functional Safety (ISO 26262) concepts and analyses
 - Unique Frequency (F) and Monitoring (M) Rating Tables



New PFMEA Severity Table

Process General Evaluation Criteria Severity (S)							
Potential Failure Effects rated according to the criteria below.							
s	Effect	Impact to Your Plant	Impact to Ship-to Plant (when known)	Impact to End User (when known)	Corporate or Product Line Examples		

AIAG 4th Edition

Issue with Severity 10/9

"Without warning" – "with warning"



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10 – Safe operation defects

9 – Noncompliance with regulations

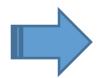


New PFMEA Occurrence Table

Occurrence Potential (O) for the Process							
Controls qual occurre FMEA	Potential Failure Causes rated according to the criteria below. Consider Prevention Controls when determining the best Occurrence estimate. Occurrence is a predictive qualitative rating made at the time of evaluation and may not reflect the actual occurrence. The occurrence rating number is a relative rating within the scope of the FMEA (process being evaluated). For Prevention Controls with multiple Occurrence Ratings, use the rating that best reflects the robustness of the control.						
О	Prediction of Failure Cause Occurring	Type of Control	Prevention Controls	Corporate or Product Line Examples			

AIAG 4th Edition

Rating based on defects/thousand, set for high volume production rates



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Rating based on robustness of prevention controls, can be applied to any production rate



New PFMEA Detection Table

Detection Potential (D) for the Validation of the Process Design						
Detection Controls rated according to the Detection Method Maturity and Opportunity for Detection.						
D	Ability to Detect	Detection Method Maturity	Opportunity for Detection	Corporate or Product Line Examples		

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Rating based on "Opportunity for Detection" and "Likelihood of Detection" by Process Controls

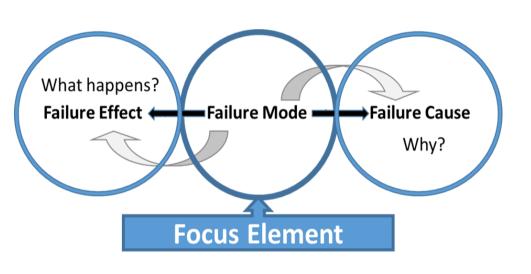


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Rating based on "Maturity of Detection Method" and "Opportunity of Detection" More stringent ratings, requires control of rejected product to prevent outflow



Failure Analysis (Step 4) – Process FMEA



Failure Chain Model

Addition of 4M

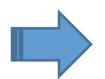
- For each Failure Mode (FM)
 consider these categories as
 sources of Failure Cause (FC)
 - Man
 - Machine
 - Material
 - Environ<u>M</u>ent



Action Priority (AP) Table

Action Priority (AP) for DFMEA and PFMEA							
Action Priority is based on combinations of Severity, Occurrence, and Detection ratings in order to prioritize actions for risk reduction.							Blank until filled in by user
Effect	S	Prediction of Failure Cause Occurring	0	Ability to Detect	D	ACTION PRIORITY (AP)	Comments

AIAG 4^{th} Edition RPN = S x O x D All three weighted equally



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S, O, D considered at the same time, while weighting Severity highest, then Occurrence, then Detection Determines Priority of Action = H, M, L



Adoption / Deployment Timing

- Expect "rolling change" not immediate changeover
 - No expectation for "rework" of existing FMEAs
 - Handbook recommends utilizing for updates to "Family" FMEAs
- Possible sequence for transition
 - 1. Allow several months for OEM and supplier training
 - 2. Acceptance of new methods, rating tables, forms on supplier FMEAs
 - 3. Then shift to requirement for new products or processes
- Expect timing for adoption / deployment from OEM's to be communicated at AIAG Quality Summit in October, 2019



