

Production Part Approval Process

Supplier Information

Production Part Approval Process Navigation

- » What's PPAP?
 - meaning
 - origin

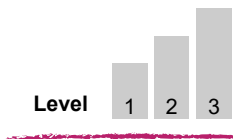


- » Benefits & Targets
 - Improve quality cooperation
 - Zero failure target Increase customer satisfaction
 - confidence in future product Risk reduction **Benefits** Part view + Process view design integrity
 - Reduce warranty charges smooth SOP managing supplier changes
 - standardization **Prevents costs for poor quality**
 - Global approach

- » What does it mean for you as the supplier?



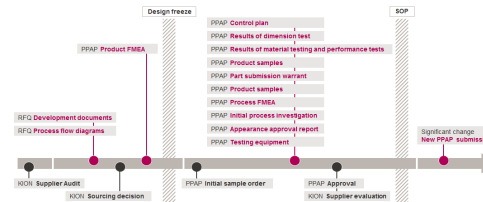
» PPAP Level



» PPAP Content

12 different requirements including example

» PPAP Timeline



» Additional information

- KION standard
- PPAP website
- contact

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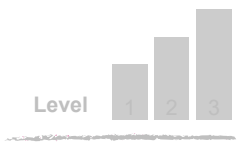
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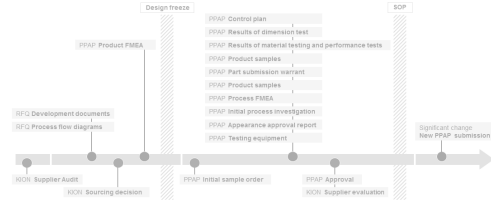
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Production Part Approval Process

What's PPAP?

P

PRODUCTION

P

PART

A

APPROVAL

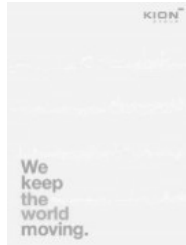
P

PROCESS

- Standard used to reduce risks prior to product release
- Team oriented approach using well established tools and techniques
- Developed by Ford, Chrysler and General Motor in 1993

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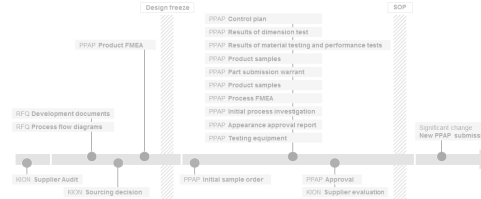
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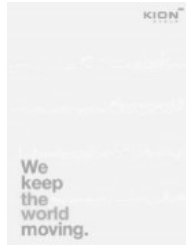
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Production Part Approval Process Benefits & Targets

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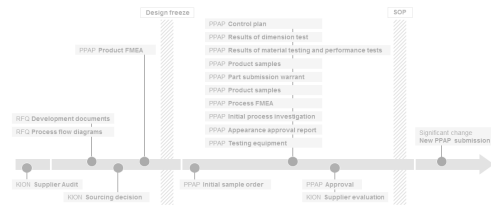
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Production Part Approval Process

What does it mean for you as a supplier?

What is the past?

- Initial sample inspection
- Part/ product view
- Different plants with own quality organization
- Different initial sampling procedures
- Huge number of suppliers in KION
- Local sourcing approach

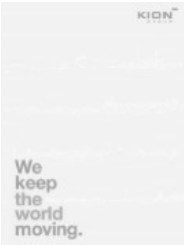
What is it like now?

- Process view added
- A stable process guarantees conform parts
- KION quality organization
- Standardized processes
- No surprise for supplier to work with all plants
- Global suppliers delivering to different KION plants

➔ Win-Win for supplier and KION

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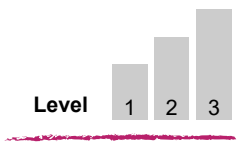
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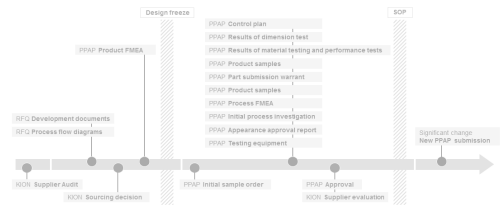
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Production Part Approval Process

PPAP Level

Level	Requirements
1	PPAP with sample parts and limited data submitted 
2	PPAP with sample parts and extensive data submitted 
3	PPAP with sample parts and complete data submitted 

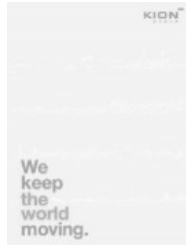
KION R&D defines the submission level depending on the following criteria :

- Complexity of part
- Complexity of process
- Safety relevance of the part
- Traceability

First angle projection 	①	②	③	④	⑤												
General tolerance (GT) in mm Size range	Index	Alteration	Change no.	Fit	Toler.												
<table border="1"> <tr> <td>≤ 30</td> <td>> 30 ≤ 120</td> <td>> 120 ≤ 400</td> <td>> 400 ≤ 1000</td> </tr> </table>	≤ 30	> 30 ≤ 120	> 120 ≤ 400	> 400 ≤ 1000	PPAP ②②	Material:	⑧	Weight in kg	⑭								
≤ 30	> 30 ≤ 120	> 120 ≤ 400	> 400 ≤ 1000														
GT fine	Inspection dim. () Aux dim. ()	Blank no.:	⑨	Title:	⑩												
<table border="1"> <tr> <td>L</td> <td>0,2</td> <td>0,5</td> <td>0,8</td> <td>2</td> <td>4</td> </tr> <tr> <td>≤</td> <td>0,1</td> <td>0,2</td> <td>0,4</td> <td>1</td> <td>2</td> </tr> </table>	L	0,2	0,5	0,8	2	4	≤	0,1	0,2	0,4	1	2	Drawn	⑥	⑦	⑪	⑮
L	0,2	0,5	0,8	2	4												
≤	0,1	0,2	0,4	1	2												
Lengths (L) and angle (∠) = ±GT	Auth'd	M. check	⑫	⑬	⑯												
Tolerance Symbols ISO 1101	⑬	⑭	⑮	⑯	⑰												
<ul style="list-style-type: none"> ○ roundness = 1/2σ-Tol. -□ straightness/flatness = AT ⊙ concentricity/run out = AT ≡ symmetry = AT // parallelism = AT ⊕ position = AT 	⑰	⑱	⑲	⑲	⑲												
Languages: ⑳	Confidential document Refer to protection notice ISO 16016	Repl. ㉑	Orig. ㉒	⑲	⑲												

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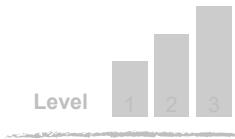
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» **What does it mean for you as the supplier?**

past → today

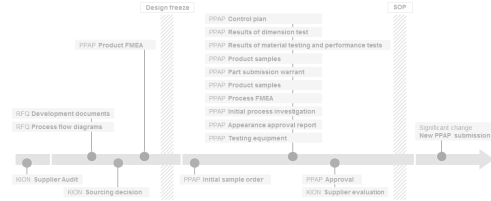
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Production Part Approval Process Requirements

	Level 1	Level 2	Level 3
1. Development documents	S	S	S
2. Product FMEA			S
3. Process flow diagrams		S	S
4. Process FMEA		R	S
5. Production control plan	R	S	S
6. Results of dimension tests	S	S	S
7. Results of material testing and performance tests	S	S	S
8. Initial process investigations		S	S
9. Appearance approval report		S	S
10. Product samples	S	S	S
11. Testing equipment			R
12. Part submission warrant	S	S	S

S – to be submitted to KION
R – to be available on request

Automotive industry: 18 criterias
KION: 12 criterias

Production Part Approval Process Requirements



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1. Development documents	S	S	S
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10. Product samples	S	S	S
11. Testing equipment			R
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Production Part Approval Process Development documents

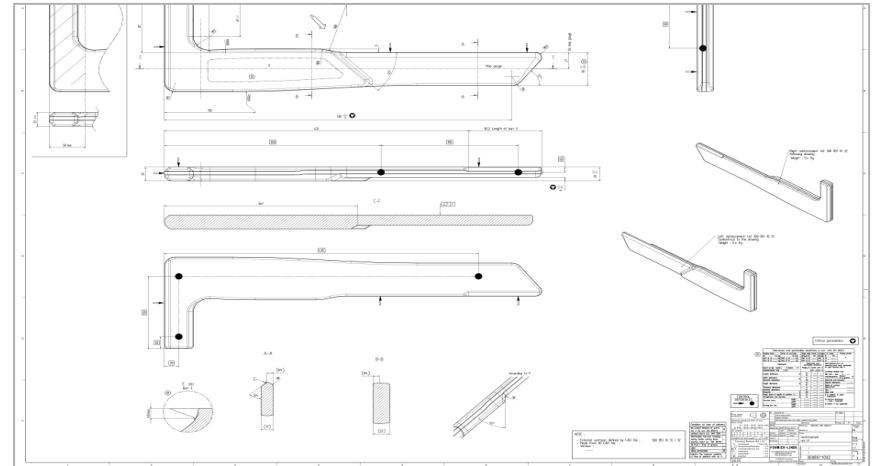


Description

Includes:

- Component drawings
- Assembly drawings
- Bill of Material
- Material specifications
- Functional/ performance specifications
- Further documents

Example



- ➔ Ensures manufacturer has the complete design record at the correct revision levels
- ➔ To be submitted with the quotation

Production Part Approval Process

Production control plan



Description

- Define measuring and testing methods including sample size/ frequency, testing equipment and documentation
- Includes reaction plan in case characteristic or process parameter is found to be out of control
- Inputs come from the supplier knowhow (Process FMEA , experiments)

KION Form

KION GROUP		Produktionslenkungsplan Control Plan						Seite 13 von 77					
Produktionslenkungsplan-Nr. Control Plan Number		1001		Table-Nr./letzte Änderungsstand Part Number/ Latest Change Level		2002970101 / 01		Anspruchspartner/ Teil Key Contact/ Piece		H. Müller / 0021 - 994708			
Datum (Erstellung) Date (Cre.)		01.08.2016		Verfahrensbezeichnung Part Name/ Description		Support		Projektteam Core Team		Maier, Müller, Schulte			
Datum (Änderung) Date (Rev.)		10.08.2016		Lieferant/ Werk Supplier/ Plant		Wiesbaden		Lieferant/ Werk/ Lieferant Datum Supplier/ Plant Approval Date		01.09.2016			
				Lieferanten-Nr. Supplier Code		25412		Andere Freigabe (RfH gefordert) Other Approval (If req'd)					
Teil-/ Prozess- schlüssel-Nr. Part/Process Number	Prozessschritt/ Arbeitsgang/ Beschreibung Process Name/ Operation Description	Maschine, Vorrichtung/ Werkzeuge zur Herstellung Machine, Device, Jig/ Tools for Mfg.	Methoden Characteristics			Methoden Methods							
			Nr. No.	Produkt Product	Prozess Process	Besondere Merkmale/ Klassifiz. Special Char. Class.	Produkt/ Prozess Spezifikation/ Genehmigte Product/ Process Specification/ I Tolerance	Werkzeugtech. Evaluation Measurement Technique	Meßsystem- analyse Measurement System Analysis Studies	Stichproben/ Samples Sampling Size	Intervall/ Frequ. Interval/ Freq.	Lenkungs- methode Control Method	Reaktionsplan Reaction Plan
10	Turning operation	5240 Turning machine GMX	13	external thread M12		no	20 +/-1	vernier caliper	Model 1	3	per 1 hour	worker self- inspection	Instruction 4- NO Parts: 1. Measurement of produced parts to the first located part 2. Change of the machine parameter 3. Messung erstes Teil mit Dreißblech

→ Ensure that the production process is under control

Production Part Approval Process

Results of dimension test



Description

- Evidence that **the complete** dimensional check has been performed
- Record results for all specified requirements out of every tool, cavity, production cell or line
- PPAP samples used for dimensional check must be identified

Example

N°	INSTRUMENT	UNIT	MEASUR	DIFF	MEASUR	DIFF	MEASUR	DIFF	MEASUR	DIFF	MEASUR	DIFF	MEASUR	DIFF	METHOD OF CONTROL / OBSERVATION	REWORK	REWORK	REWORK	REWORK
1	() 100	-2	200,06		99,96		99,93		99,93		99,87		99,87		pe	99,90		0,09	
2	() 152	-2	151,67		151,70		151,90		151,17		151,70		151,70		pe	151,83		0,13	
3	() 79	-2	78,46		78,38		78,46		78,27		78,41		78,40		pe	78,40		0,08	
4	() 14	-1	14,07		14,17		14,20		14,08		14,07		14,11		pe	14,11		0,08	
5	() 6	-1	5,95		5,93		6,06		6,05		5,94		5,99		pe	5,99		0,08	
6	MFC	0	0	0	0,3										pe				
7	ØA	0	0	0	0,3										pe				
8	() 23	-1	24,82		24,78		24,77		24,82		24,78		24,78		pe	24,81		0,06	
9	MFC	0	0	0	0,2										pe				
10	sondes	/	/	/	/										visual				
11	sondes	/	/	/	/										visual				
12	precision	/	/	/	/										visual				
13	Ø1	-0,3	79,93		79,94		79,94		79,89		79,91		79,90		pe	79,90		0,04	
14	Ø2	-0,3	0,3												pe				
15	Ra2	0	-0,3	0,3											pe				
16	Ø1A	0	0	0	0,07		0,10		0,08		0,07		0,11		colonne	ØZPT		ØZPT	
17	Ø1A	0	0	0	0,1										colonne	ØZPT		ØZPT	
18	ZØ8	0	0	0	0,3										pe				
19	Ø6	0	-0,2	0,2											pe				

➔ If dimensions are not conformed to the specification, the manufacturer informs KION R&D and Quality prior to PPAP submission

Production Part Approval Process

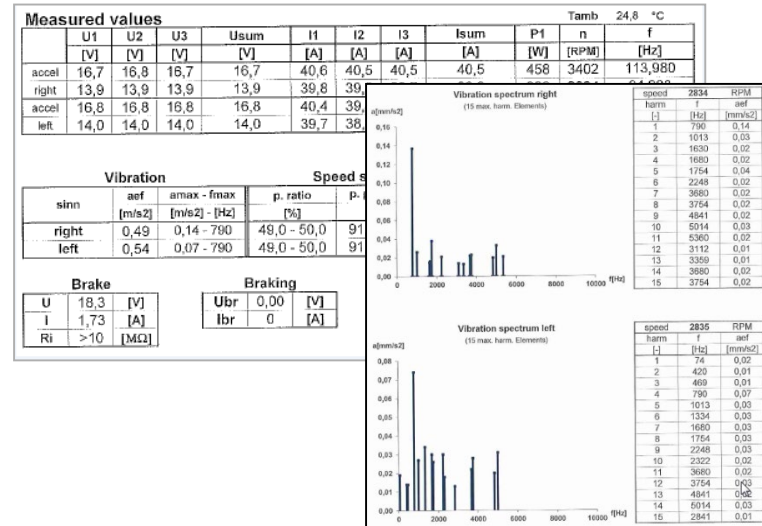
Results of material testing and performance tests



Description

- Evidence that material, performance or functional tests according to requirement (development documents or production control plan) have been performed
- Record results including test report number, date, laboratory that performed the test

Example



➔ Responsibilities for performance tests are fixed in cooperation with the KION R&D

Production Part Approval Process

Product samples



Description

- Quantity of samples defined by KION
- Product sample parts to be identified with specific label
- The product samples delivered are the parts that have been measured for the dimensional results

Example

An example of a yellow label for the Production Part Approval Process (PPAP). The label contains the following text:

Part-Number: _____ ind.: _____ ¶
Quantity: _____ ¶
Order-Nº: _____ ¶

**Production Part ¶
Approval Process ¶
(PPAP) ¶**

NB: The initial samples shall not be packed with pre-serial or serial parts. ¶

Production Part Approval Process

Part submission warrant (PSW)



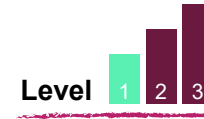
Description

- With the part submission warrant, the supplier confirms the design and the processes to manufacture the parts (including quantity of tools cavities, production cells or lines)
- Declaration that the results meet the drawing and specification requirement

KION Form

Part Submission Warrant (PSW)				KION GROUP	
Part Name	Support	Cust. Part Number	12345678		
Shown on Drawing No.	12345678	Org. Part Number	98764 - 01		
Engineering Change Level	3	Dated	21. Mrz 16		
Additional Engineering Changes	no	Dated	-		
Safety and/or Government Regulation	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No	Purchase Order No.	490000001		
Weight (kg)	1.2000	Dated	08. Aug 16		
Checking Aid No.	2432	Checking Aid Engineering Change Level	01		
ORGANIZATION MANUFACTURING INFORMATION			CUSTOMER SUBMITTAL INFORMATION		
Herr Mustermann / 1254			Meyer GmbH / München		
Organization Name & Supplier/Vendor Code			Customer Name/Division		
Max-Mustermann-Str.			Frau Maler		
Street Address			Buyer/Buyer Code		
Hamburg			Stapler xyz		
City	Region	Postal Code	Country	Application	
REASON FOR SUBMISSION (Check at least one)					
<input checked="" type="checkbox"/> Initial Submission			<input type="checkbox"/> Change to Optional Construction or Material		
<input type="checkbox"/> Engineering Change(s)			<input type="checkbox"/> Supplier or Material Source Change		

Production Part Approval Process Requirements



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1. Development documents	S	S	S
2. Product FMEA			S
3. Process flow diagrams		S	S
4. Process FMEA		R	S
5. Production control plan	R	S	S
6. Results of dimension tests	S	S	S
7. Results of material testing and performance tests	S	S	S
8. Initial process investigations		S	S
9. Appearance approval report		S	S
10. Product samples	S	S	S
11. Testing equipment			R
12. Part submission warrant	S	S	S

S – to be submitted to KION
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Production Part Approval Process Process Flow Diagrams



Description

- Designation and clear description of every single process step required to produce the referenced product

KION Form

KION		Prozessflussdiagramm Process Flow Diagram			
Teile-Nr / Mittel Änderungsstand Part Number / Last Change Level		12345678 / 03	Teilebezeichnung Part Name / Description		Master
Anspruchspartner / Tel. Key Contact / Phone		Herr Max Mustermann / 09221100xx	Lieferanten-Nr. Supplier Code		1234
Datum (Erstellung) Date (Orig.)		01.08.2016			
Lieferant/ Werk Supplier Part		Aachtfarburg			
Legende: Legend:					
Arbeitsvorgang Step	Arbeitsvorgang oder Ereignis Operation or Event			Beschreibung des Arbeitsvorgangs oder des Ereignisses Description of Operation or Event	
10					Goods receipt storage
20			X	X	Transport production turning machine

- ➔ Provide linkage to Process FMEA and Control Plan
- ➔ Has to be submitted with the quotation

Production Part Approval Process Process FMEA



Description

- Process Failure Mode and Effects Analysis:
 - determine the ways in which the Process Steps can go wrong including effects and causes
 - List the current controls for each potential cause
 - Evaluate the risk (RPN)
 - Define recommended actions
- Inputs come from the process flow diagram

KION Form

Failure Mode and Effects Analysis

Process name		Owner	Creation date	Item origin	ISO analysis
Process: Support - Turning operation		Walter	01.06.2016	Counterbalance truck vpt	100
Project		Walter		Referenced: 07.08.2016	100
Support		Walter		07.08.2016	100
Project change		Walter		07.08.2016	100
Milestone		Manufacturing	Manufacturing - Dep 1	07.08.2016	100
Walter		Manufacturing	Manufacturing - Dep 1	07.08.2016	100
Walter		Manufacturing	Manufacturing - Dep 1	07.08.2016	100
Walter		Manufacturing	Manufacturing - Dep 1	07.08.2016	100
Walter		Manufacturing	Manufacturing - Dep 1	07.08.2016	100

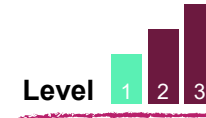
No.	Component / Function / Process	Failure mode	Failure effect of failure	Failure failure causes	Current process control	S/N			Recommended Action	Target date	Action taken			S/N		
						Sy	Severity	Occurrence			Sy	Severity	Occurrence			
1	Turning operation 1	Dimension is below tolerance	Press fit between holder and pin not possible	Incorrectly adjusted tool	Visual inspection	5	5	4	100	Invest tool presetter	Walter	Invest tool presetter	1	5	4	20
		Dimension is above tolerance	Support not applicable in the drill hole	M incorrectly measurement of the tooling	Visual inspection in the turning machine	5	5	4	100	Invest tool presetter in the turning machine	Walter	M invest tool presetter in the turning machine	1	5	4	20
						5	5	4	100				1	5	4	20
						5	5	4	100				1	5	4	20
						5	5	4	100				1	5	4	20
						5	5	4	100				1	5	4	20
						5	5	4	100				1	5	4	20

Severity X Occurrence X Detection = RPN
5 X 4 X 2 = 40

➔ Carry over process controls and recommended actions into the control plan

Production Part Approval Process

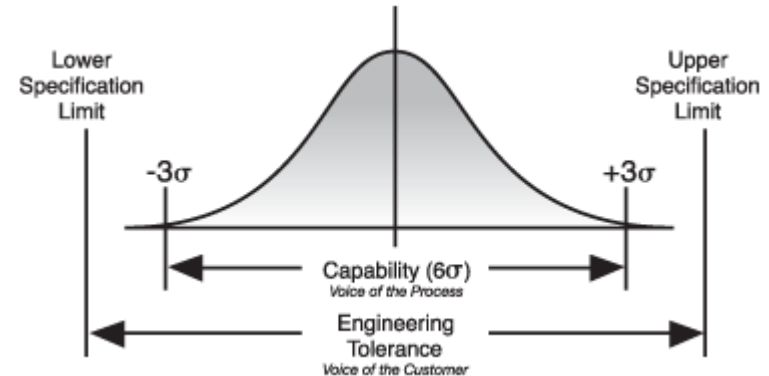
Initial process investigations



Description

- determine how well the production process can manufacture products that meets the design requirements (stable and capable)
- Short-term investigations for special characteristics according to the drawing
- The number of samples for the initial investigations depends on the supply quantity over one year
- Supplier agrees with KION index to be used to determine the initial process capability

Example



→ May be replaced by long-term results from identical or similar processes (with KION agreement)

Production Part Approval Process

Appearance approval report



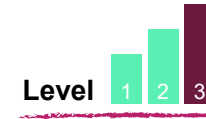
Description

- Only suitable for products with color, grain or surface appearance requirement
- Appearance approval for families or individual parts is acceptable

KION Form

APPEARANCE APPROVAL REPORT															KION GROUP						
PART NUMBER					DRAWING NUMBER					APPLICATION											
PART NAME					BUYER CODE			E/C LEVEL			DATE										
SUPPLIER NAME					MANUFACTURING LOCATION					SUPPLIER CODE											
REASON FOR SUBMISSION					<input type="checkbox"/> PART SUBMISSION WARRANT			<input type="checkbox"/> SPECIAL SAMPLE			<input type="checkbox"/> RE-SUBMISSION			OTHER							
					<input type="checkbox"/> PRE TEXTURE			<input type="checkbox"/> FIRST PRODUCTION SHIPMENT			<input type="checkbox"/> ENGINEERING CHANGE										
APPEARANCE EVALUATION																					
ORGANIZATION SOURCING AND TEXTURE INFORMATION												PRE-TEXTURE EVALUATION		AUTHORIZED CUSTOMER REPRESENTATIVE SIGNATURE AND DATE							
												CORRECT AND PROCEED									
												CORRECT AND RESUBMIT									
												APPROVED TO ETCH/TOOL/EDM									
COLOR EVALUATION																					
COLOR SUFFIX	TRISTIMULUS DATA					MASTER NUMBER	MASTER DATE	MATERIAL TYPE	MATERIAL SOURCE	HUE				VALUE	CHROMA	GLOSS		METALLIC BRILLIANCE		COLOR SHIPPING SUFFIX	PART DISPOSITION
	DL*	Da*	Dd*	DE*	CMC					RED	YEL	GRN	BLU			LIGHT	DARK	GRAY	CLEAN		

Production Part Approval Process Requirements

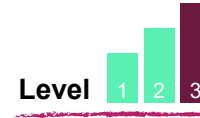


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2. Product FMEA			S
3. Process flow diagrams		S	S
4. Process FMEA		R	S
5. Production control plan	R	S	S
6. Results of dimension tests	S	S	S
7. Results of material testing and performance tests	S	S	S
8. Initial process investigations		S	S
9. Appearance approval report		S	S
10. Product samples	S	S	S
11. Testing equipment			R
12. Part submission warrant	S	S	S

S – to be submitted to KION
R – to be available on request

Production Part Approval Process

Product FMEA = Design FMEA



Description

- If supplier is responsible for the design
- **Design Failure Mode and Effects Analysis:**
 - Provide potential cause
 - Evaluate Risk Priority Number (RPN) of failures
 - Define corrective and preventive actions (Re-design, DVP&R...)
- Consider functions from product or system
- An individual Product FMEA may be applied to a family of similar individual parts

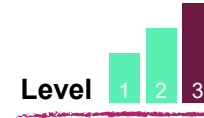
KION Form

Severity	X	Occurrence	X	Detection	=	RPN
5		4		2		40

➔ To be submitted before design freeze

Production Part Approval Process

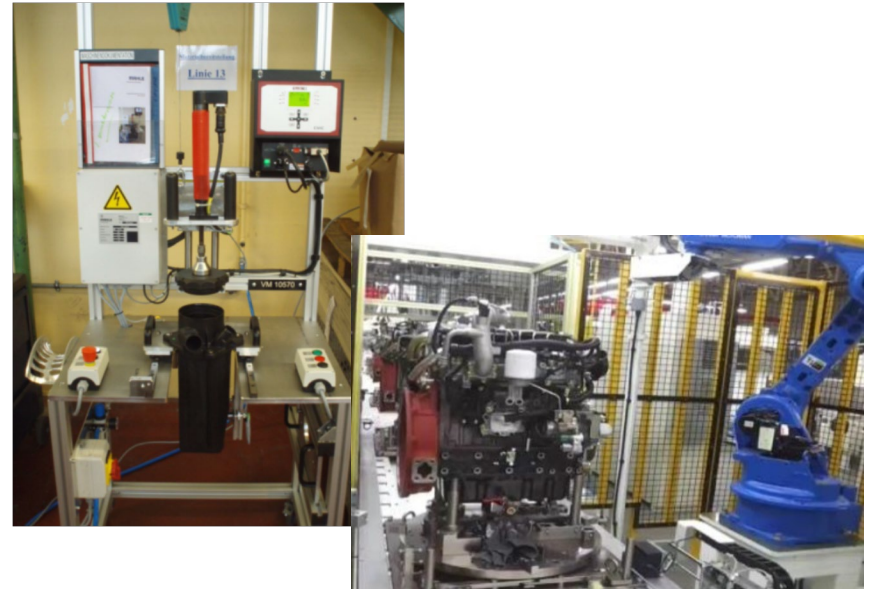
Testing equipment



Description

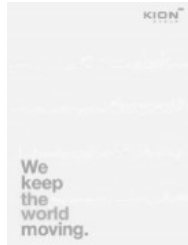
- If required by KION, the supplier shall submit the part-specific testing equipment for assemblies or components as part of the PPAP submission

Example



Production Part Approval Process Navigation

- » What's PPAP?
 - meaning
 - origin

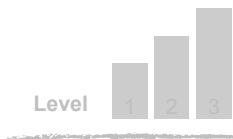


- » **Benefits & Targets**
 - Improve quality cooperation
 - Zero failure target Increase customer satisfaction
 - confidence in future product **Benefits** Part view + Process view
 - Risk reduction design integrity
 - Reduce warranty charges smooth SOP managing supplier changes
 - standardization **Prevents costs for poor quality**
 - Global approach

- » What does it mean for you as the supplier?



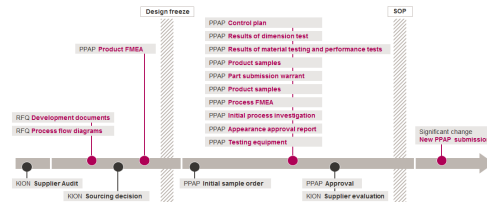
- » **PPAP Level**



- » **PPAP Content**

12 different requirements including example

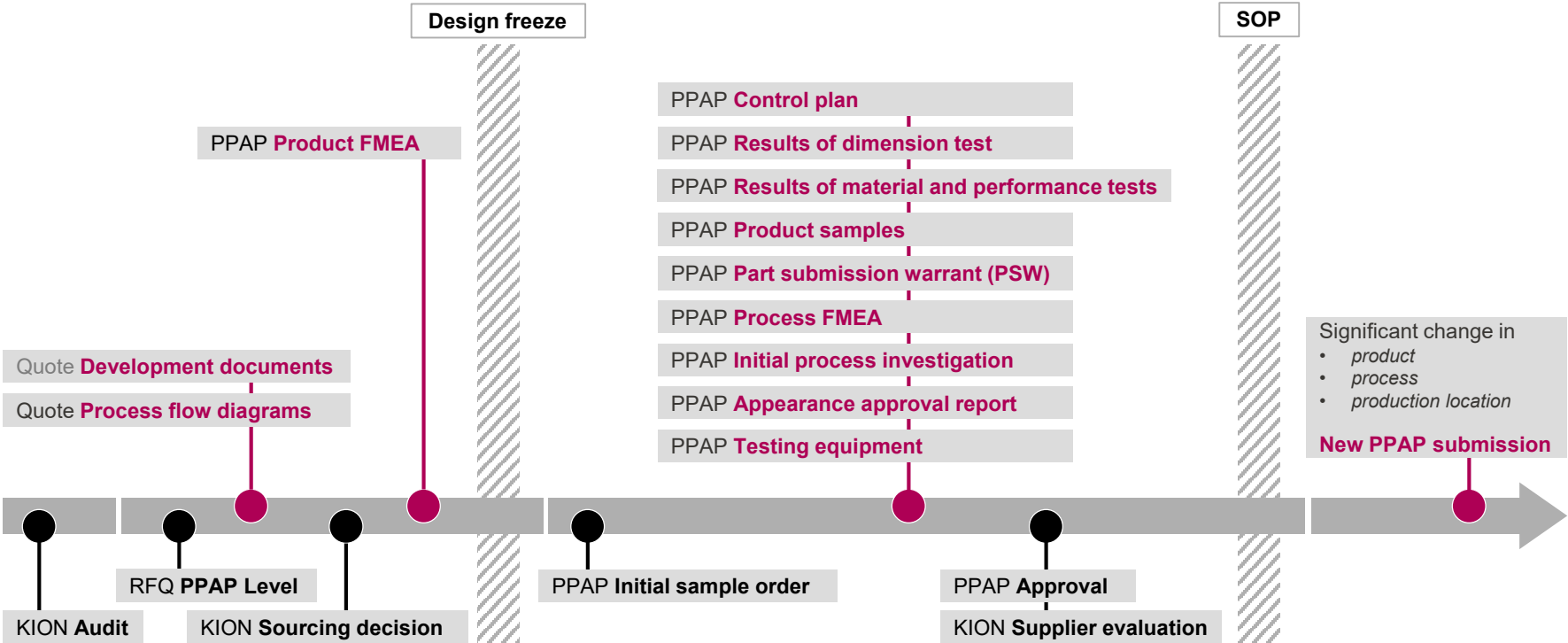
- » **PPAP Timeline**



- » **Additional information**

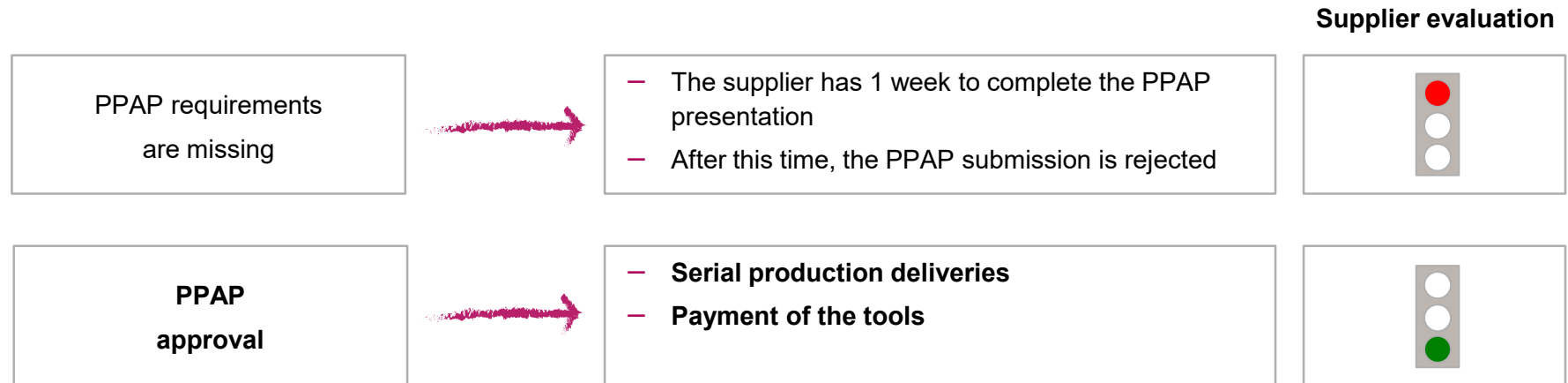
- KION standard
- PPAP website
- contact

Production Part Approval Process Timeline



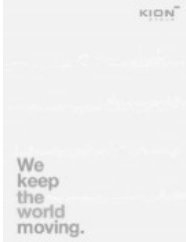
Production Part Approval Process

PPAP status consequences



Production Part Approval Process Navigation

- » What's PPAP?
 - meaning
 - origin



» **Benefits & Targets**

Improve quality cooperation
Zero failure target Increase customer satisfaction

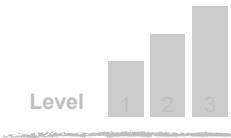
confidence in future product
Risk reduction **Benefits** Part view + Process view
design integrity

Reduce warranty charges smooth SOP managing supplier changes
standardization **Prevents costs for poor quality**
Global approach

» What does it mean for you as the supplier?



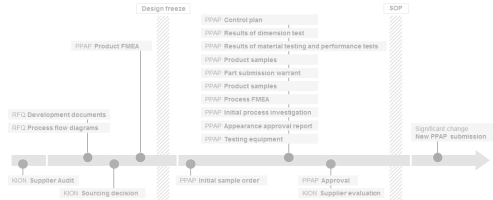
» **PPAP Level**



» **PPAP Content**

12 different requirements including example

» **PPAP Timeline**



» **Additional information**

- KION standard
- PPAP website
- contact

Production Part Approval Process Additional Information

- » • **KION Standard**
– WN 50020

Works Standard		WN 50 020
KION GROUP		
PPAP - Production Part Approval Process		
1 AREA OF APPLICATION AND PURPOSE		
This area describes the general, basic requirements for the approval of vendor parts for production. It is based on the requirements for external control as IATF 16949. It also describes the specific requirements for the necessary process and supply products that meet the quality requirements. The standard applies to the KION Group and all suppliers.		
2 TERMS		
Representative of KION Quality – Contact in the Supplier Quality Department COP – Process Capability Index PPAP – Process Part Approval Process COP – Process Control Plan COP – Control Plan COP – Control Plan COP – Control Plan COP – Control Plan COP – Control Plan		
3 GENERAL		
The supplier shall submit a new approval PPAP before the first series delivery in the following situations (provided that the representative of KION Quality has not raised the requirements):		
<ul style="list-style-type: none"> - A new individual part, a new component or a new product - Correction of a part that has already been submitted - An engineering change (new individual part number) - Any situation in which a new approval PPAP shall be submitted in accordance with Section 4. 		
4 REQUIREMENTS FOR THE PPAP PROCESS		
4.1 Representative production run		
In the case of production parts, the products for the PPAP (initial samples) shall be taken from a representative production run. This representative production run is defined using a specific production quantity of consecutive individual parts, which shall be agreed between the supplier and the representative of KION Quality.		
This representative production run shall be carried out at the production site using the appropriate production tools, gauges, processes and supplies, and shall include the usual production conditions, other series conditions.		
Individual parts from each individual production process (e.g. from every other assembly line or production cell) from each position in the case of multiple process lines or process cells shall be released and/or tested.		
4.2 PPAP Requirements		
The supplier shall fulfil all the PPAP requirements specified and listed below (see 4.2.1 to 4.2.12). Components and individual parts shall fulfil all the requirements contained in the technical development documents and the specifications from KION (including parts and quantity requirements).		
4.2.1 Development documents (drawings, CAD files)		
The supplier shall have the development documents (drawings, specifications and/or documents) for the specific production/individual part, components or units. This shall be verifiable with the supplier.		
Comment 1: For each deliverable product, each individual part or each component, only one development document is released. Changes to KION (e.g. any change of development documents, approvals of new or replacement documents, the development documents may refer to other documents, which will become an integral part of the development documents.		

- » • **Website**
– KION PPAP website

<https://www.kiongroup.com/en/Service/KION-PPAP/>

- » • **Contact**
– Email/ Phone

Please contact the
Quality@Supplier representative
for specific questions

KION[®]

G R O U P