Quality is Everyone's Responsibility Digital Workflows for Quality Management Excellence

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Designed by Cambrian Lab



Quality is everyone's responsibility - W. Edwards Deming

Agenda

- Introductions
- □ Cross-functional Quality Management and Challenges
- □ Workflow Based Approach to Cross-functional Quality Management
- □ Case Study
- Demo
- Q&A



About Cambrian Lab

Team

Supply Chain, New Product Introduction, and Technology Experts from SAP, Samsung, Siemens, GM, Ford, Applied Materials

Industries

Automotive, High-tech, Semiconductor Equipment, Medical Devices, Consumer Products (From Fortune 100 to Start-ups)

Locations

SF Bay Area, Detroit, Boston, Houston

Expertise

- New Product Development/Introduction, Supplier Development, and Quality Management
- Enterprise and Supply Chain Technology (ERP, SCM, Manufacturing, CRM, Sourcing, Finance)



Panelists Today

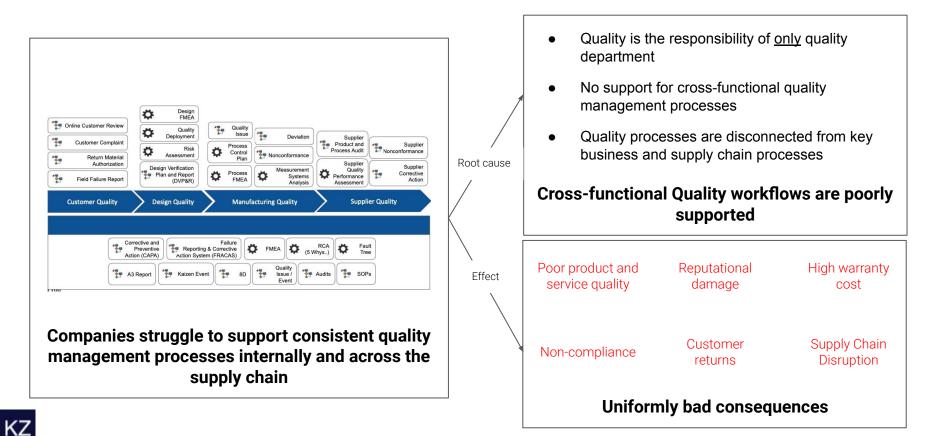
Kris Gorrepati

- 20+ years experience in New Product Development and Introduction and Supply Chain Manufacturing
- SAP, Samsung, Ford, Caterpillar
- Auto, High-tech, Software
- Michigan Tech (Mech Engg.), UCLA

Manish Mathur

- 20+ years experience in PLM, Business Analytics, Software Engineering
- GM, Ford, Siemens PLM, Cap Gemini
- Auto, High-tech, Software
- Michigan Tech (Elec Engg), Walsh College

Making Quality Everyone's Responsibility is a Challenge



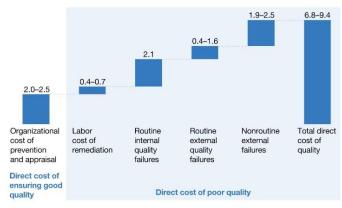
Cost of Poor Quality >>>> Cost of Quality

Cost of Quality Cost of Good Quality Cost of Poor Quality						
Prevention costs	Appraisal costs	Internal failure costs	External failure costs			
Planning Process Control Quality Audits Maintenance Supplier Evaluation Training Design Review Risk Assessment FMEA	 Inspection FAT Document Review Quality Audits Calibration Test Materials Test product 	 Scrap Rework Missing documents Problem solving Retest Redesign Downgrading Variation Downtime 	 Warranty charges Complaints Returned Materials Late Delivery Penalties Rework After Installation Lost Opportunities 			

For the first nine months of 2020, Ford's warranty costs totaled \$3.87 billion, while GM's were \$1.68 billion, according to regulatory filings.

The direct cost of quality in the medical device industry is \$26 billion to \$36 billion annually.





¹Estimated annual sales of \$380 billion.

McKinsey&Company | Source: Health Research International; McKinsey analysis

Why Making "Quality is Everyone's Responsibility" Remains a Challenge

To "Do What is Documented" as QMS is Hard

Manual

QMS via Email

Actual practice varies from what is expected

Requires extraordinary effort from people

Involving Supply Chain is Difficult

Suppliers not in the loop

No visibility into Supplier quality practices

Cross-functional teams and systems are disconnected

Missed handoffs

Quality events fall through the cracks

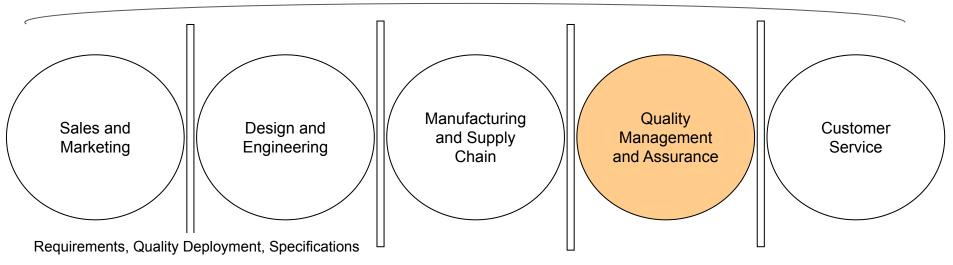
Institutional Knowledge lost

No guarantee that corrective/preventive actions are implemented effectively and in a timely manner

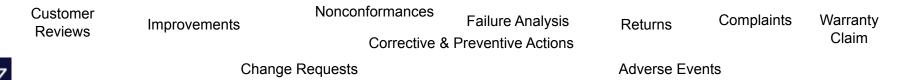
No learning from prior quality events Dropped or delayed corrective and preventive action implementation



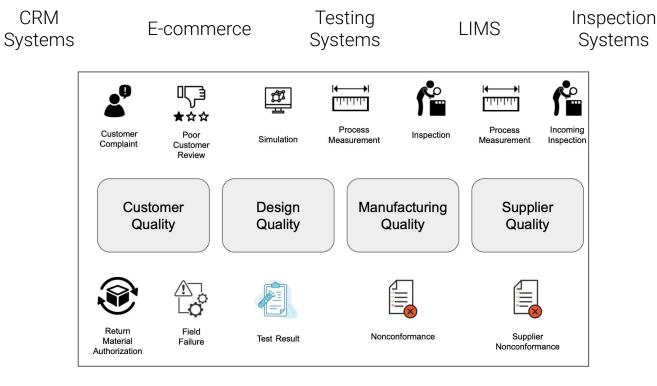
Process and Information Silos Are Obstacles to Making "Quality is Everyone's Responsibility"



FMEA, Validation, Prototyping, Product and Process Characteristics, Process Plan, Control Plans, SOPs, Training, Audits..

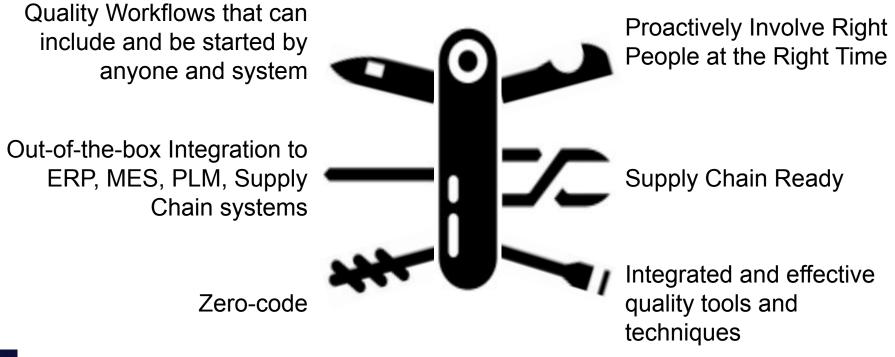


Quality Events can get Lost in Disparate and Disconnected systems

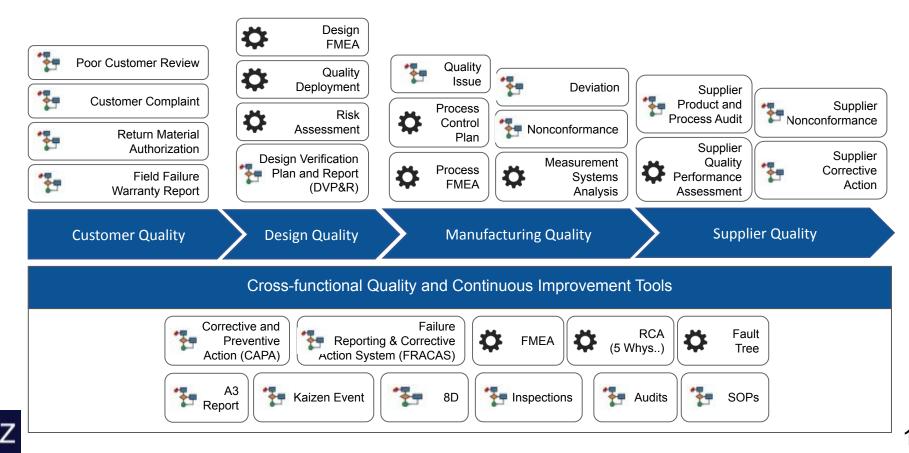




ERP Systems Supply Chain Systems The Answer - Cross-functional Workflows that Make "Quality is Everyone's Responsibility" a Reality



Quality is Everyone's Responsibility



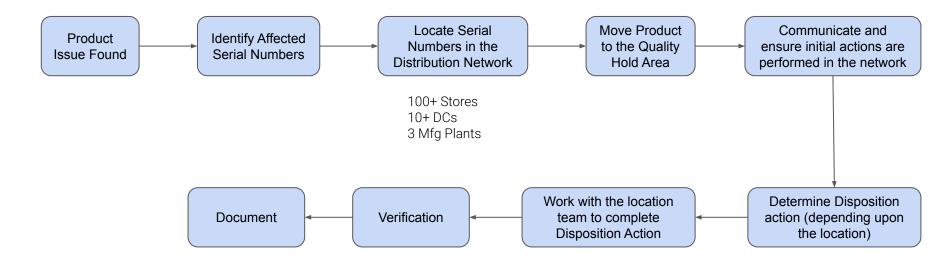
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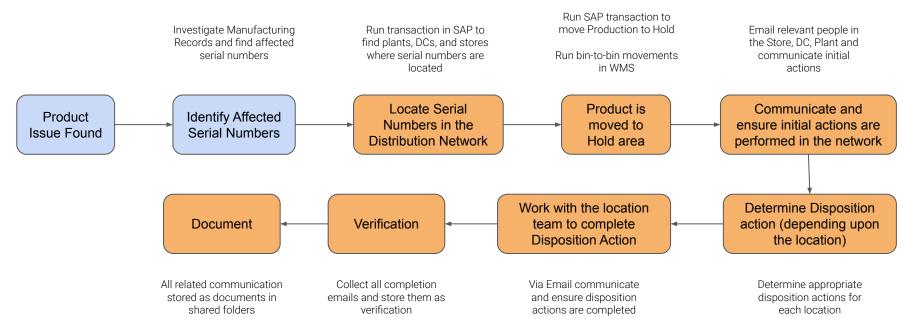


Case Study Overview

Manufacturer needs to act fast to move non-conforming product (so that it does not end up with customers) from various distribution nodes (Stores, DCs, and Manufacturing Plants) to support immediate actions, fixes and eventual disposition

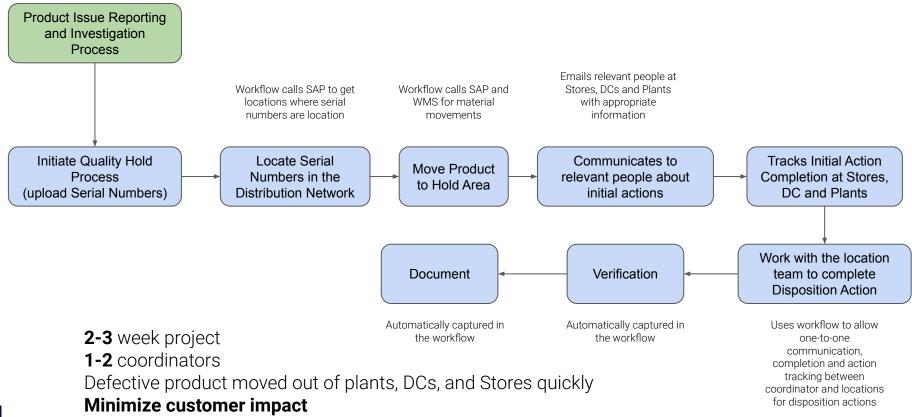


AS-IS Standard Operating Procedure



3-6 month project
10-20 people involved
High chance of defective product making it to customers
Very expensive and damaging to the brand if defective product ends up with customers

Digital Workflow Based Standard Operating Procedure



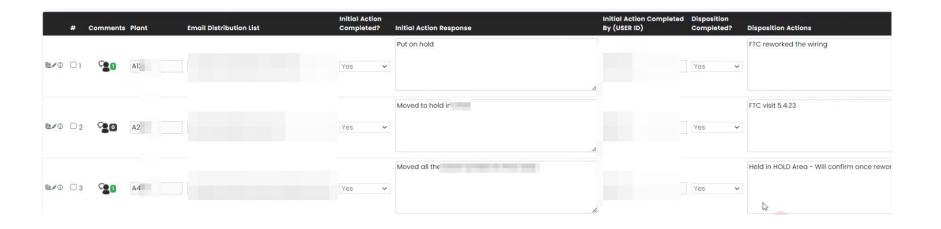
Initiate Product Hold

	Serial Number		
nfo Current Activity	Data Members Roles WorkFlow Log Graph Process Report	# Se	rial Number
urrent Activity: Wait for Plan	Action Completion (Current Status: Activity restarted) You are Acting for the Performer Performers:	01	254
mary Facility:	~	D 2	255
porting Store/DC/Facility :@	<- Select -> 🗸	🗆 3 🗍	256
erence ID:⑦		4	257
e Description:	Wiring Issue	05	258
	10021	0 6	280
al Action Instructions:	Put these items in hold Disposition Plan - FTC's will visit for rework - Please respond Back when complete	07	291
		0 8	292
rent Scope*:	Disposition 🗸	0.9	293
t Update:	2023/05/04 14::	C 10	294

Workflow connects to SAP, gets Locations of Serial Numbers, and initiates movement to Quality Hold Area in WMS

#	Material Number	Equipment Number	Description	Serial Number	Plant	Current S Loc Customer	Impacted Quantity
🗆 I				38	A12	0001	
2				8	A12	0001	
□ 3				20	Al2	0001	
□ 4				38	Al2	0001	
0 5				38	A12	0001	
□ 6				3	A2	0001	
7				4	A2	0001	
08				2	A2	0001	
9					A2	0001	
□ 10					A4	0001	
\Box 11	IN SAL	1050000410	0435-10/044-65-00 California	0000000		0001	

Workflow Identifies and Communicates with Relevant Stakeholders at Impacted Locations



Workflow Allows bidirectional communication and coordination of initial actions, fixes and eventual disposition

			12587
Initial Action Instructions:	Put these items in hold Disposition Plan - FTC's will visit for rework - Please respond Back when complete	Disposition Instructions:	
Current Scope:	Disposition ~	Creation Date:	2023/05/04 14:06:56
Last Update:	2023/05/04 14:22:31		
Plant			
Plant:	1001	Email Distribution List:	Moved all the serial numbers to HOLD area
Initial Action Completed?:	Yes 🗸	Initial Action Response:	
Initial Action Completed By (USER ID):		Initial Action Date:	2023/05/04
Impacted Quantity Summary:	Qty: 10 ▲ Qty: 15	Disposition Completed?:	Yes 🗸
Disposition Actions:	Held in HOLD Area - Will confirm once reworked	Disposition Completed By (USER ID):	
Disposition Date:	2023/05/04		
Attachments			
CHOOSE FILE No file chose	en		

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Demo

Systematically address minor and major quality issues

Learning from Failure

Involving Suppliers



Ready to Use Cross-functional Quality Workflows, Tools, and Techniques

	Launch Pad fo	or Administrator		🔅 NPI Review	APQP Program	First Article Inspec
			1	List Find New	List Find Insights New	List Find Insights New
ossfunctional Quality						List (Find) margins (New
Corrective Action Request	🚰 🛛 Failure Report	Custom Corrective Action Report	Nonconformance Report	Supplier PPAP	昌 DFMEA	Design Verification
List Find Insights New	List Find Insights New	List Find Insights New	List Find Insights	List Find Insights	List Find Insights	List Find Insights
🚰 Quality Issue				E QFD	E Requirement	E Risk Assessment
List Find Insights New				List Find Insights	List Find Insights	List Find Insights
ontinuous Improve	ment			Supplier Quality		
		lem Solving		Supplier Performance	Supplier Risk Assessment	Supplier Audit
A3 Report	Process	left solving	🚰 Kaizen Event	List Find New	List Find New	List Find Insights N
A3 Report	Process	nsights New	List Find Insights New	List Find New		List Find Insights N
	Process			Ust Find New		List Find Insights No
List Find Insights New	Process			Supplier Performance	List Find New	
List Find Insights New anufacturing Quality	Control Plan		List Find Insights New	Supplier Performance	List Find New	Supplier Profile
List Find Insights New	List Find I	nsights I New	List Find Insights New	Report List Find Insights New	List Find New	Supplier Profile
List Find Insights New anufacturing Quality To a section Report List Find Insights New	Control Plan	Insights I New	List Find Insights New	Supplier Performance Report List Find Insights New Customer Quality	List Find New	E Supplier Profile
List Find Insights New anufacturing Quality To a section Report List Find Insights New	Control Plan	Insights I New	List Find Insights New	Supplier Performance Report Ust Find Insights New Customer Quality	List Find New ************************************	E Supplier Profile List Find Insights

Medical Device NPI

Part/Assembly

List | Find | Insights

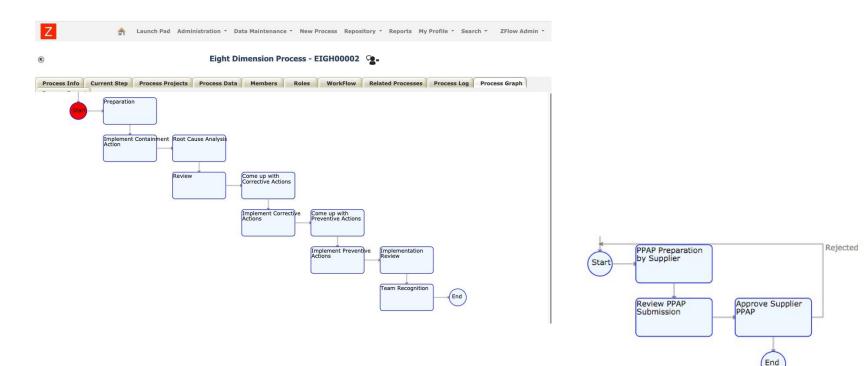
Supplier Corrective Action

Return Material

Authorization

1

Powerful Do-it-yourself Workflow Design and Execution

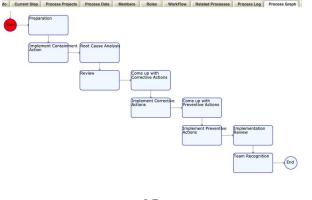


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Integrated and Effective Techniques as part of Quality Workflows

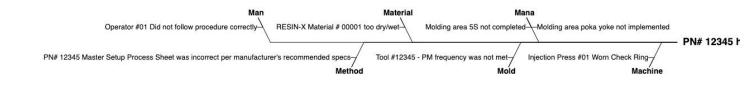
ζ.	Desi	gn FMEA - Delta Eclips	e XZ					
		E In Show		S 🔏 🗄	3			
	Show Badshe	Attachmente Attachment	Reports Cit	ose Comments Action	Items			
	gn FM		Form Identifie	Delta Eclipse XZ	Part Number:	Delta Eclipse XZ	Description:	Delta Eclipse XZ DFMEA
		Delta Eclipse XZ Maintains designed level	Status:	Work In Process	Rev.:	A	Rev.Status:	Pending
C		of low pressure	Created By:	admin@zflow.io	Creation Date:	2020/12/14	Updated By:	admin@zflow.io
С	(),	Desired vacuum level not maintained				8:37:54		
С	B ,	Contamination of test environment	Last Update:	2020/12/14 8:37:54				
С	۱ 1	Leak due to o-ring failure	Design FMEA	Function				
С	•	Continuous measurement	# 14 S	System		Syste	m Interface	
С		New o-ring	1	Vacuum Chamber				
C		Vacuum system failure		Motion Controller				
С	i ,	Continuous measurement	1000	Thermal Controller				
С		Controls movement of IC test beds	3	I hermal Controller				
С		Maintains temperature						

Eight Dimension Process - EIGH00002 @



8D

FMEA





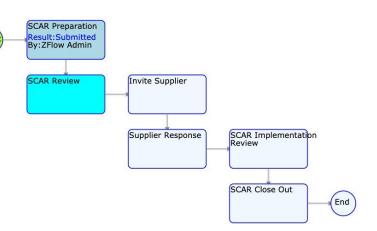
Fishbone

Cross-functional Team Collaboration

Z	Roles -	Supplier PPAP			×					
New Edit Delete Update Processes										
Role Name Description	on From Org) Privileges								
Participant		Z Admin	istration 🝷 Data M	aintenance 🔻	New Process	Repository	• Reports	My Profile	 Search * 	ZFlow Admin
) Design Engineer	No			Organ	nization Str	ucture - R	oot			
) Process Admin		Edit Organization Position Use	New Export Rename	X Delete De	elete All Expand Al	87.000	een Org Repor	t Download	(S) Back	
Durchasing Load	No	Entity		Туре	Person	Assigned			Reports	To Position
Purchasing Lead	NO	🔿 📑 Root		Organization						
		 ZZ Corp 		Company						
Supplier Representative	No	🔿 📑 Tezla		Organization						
		O O CEO		Position	zflow-ce	eo@mailinator	.com Ø			
		O VP of Finance		Position	zflow-vp	o-finance@ma	ilinator.com	đ		
		O VP of Manufactu	ring	Position						
		🔿 🛛 📑 Fremont Plant		Division						
		O Depart Manager		Position						
		🔿 📑 Giga Factory		Division						
		🔿 📑 Z Lab		Organization						

Easy and Secure Supply Chain Collaboration

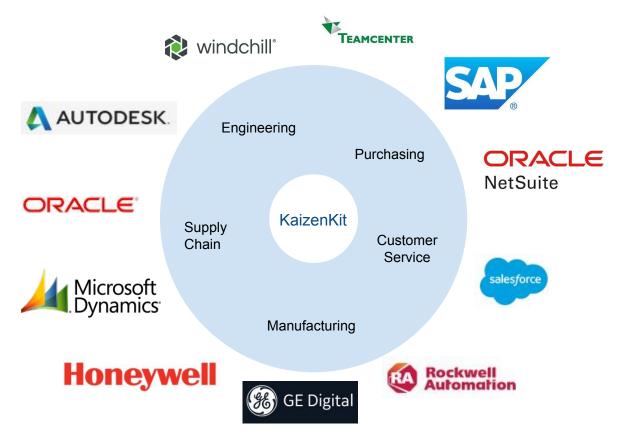
-					
	Business Name	Supplier ID	Supplier Relationship Status	Supplier Maturity Level	Supplier Category
	Avioq Inc.	SAVIOQ	Active	Advanced	Strategic
	Bosch Sensortec	SBOSCH	Qualified	Advanced	Non -strategic
	Broadcom	SBRCM	Active	Advanced	Strategic
	Carrier UTC	Carrier			
	Cirrus Logic	SCIRRUS			
	Genco Contractors	Genco			
	Generac	Generac			
	Honeywell	Honeywell			
	Huizhou Desay	SHUIZHOU		Basic	Non -strategic
	Intel Corp	SINTC	Preferred	Integrated	Strategic
	Jabil	Jabil	Preferred	Integrated	Strategic
	Lattice Semiconductor	SLATTICE	Qualified	Advanced	Non -strategic
	Lennox International	Lennox			
	MetalDyne	MetalDyne	Active	Basic	
	Murata Manufacturing	MURATA	Active	Advanced	Routine
	Northern Tool and Equipment	NorthernTE			
	NXP Semiconductor	NXPSEMI	Preferred	Advanced	Non -strategic
	Qorvo Inc	QORVO	Qualified	Advanced	Non -strategic
	Qualcomm Inc.	QCOM	Restricted	Advanced	Strategic
	Skyworks Inc.	SKYWORKS	Active	Advanced	
	TDK Corp	TDKCORP			



Quality Workflow Visibility and Metrics



Rich Capabilities for Integrating to ERP, Engineering, Manufacturing and Supply Chain Applications





Summary

- *Proven success* in many industries
- □ Easy to adopt and loved by users
- □ Easy to get started





For More Information

https://www.kaizenkit.io

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