



A TECHWELL EVENT

W16

Test Automation

Wednesday, October 17th, 2018 3:00 PM

Delivering the Goods: Harmonizing Regulated and Agile Practices

Presented by:

Griffin Jones

Brought to you by:



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Griffin Jones

Griffin Jones is an Agile consultant and coach, specializing in applying human-centered systems thinking to regulated industries. He has a technical background of over twenty-five years of testing as an individual contributor, Manager, Director, and Consultant - for medical device and pharmaceutical companies, clinical research organizations, banks, and insurance companies. Griffin has been responsible for all matters relating to the establishment and maintenance of a Quality Systems, the execution of verification and validation (testing), and the presentation and representation of results to internal and external regulatory auditors. He is currently a host of the Workshop on Regulated Software Testing (WREST). Reach Griffin at griffin.jones@congruentcompliance.com.



Delivering the Goods

***Harmonizing
Regulated and Agile Practices***





Griffin Jones

Consultant / Coach

Agile / Testing / Regulatory

This Presentation Covers ...

- **My 25+ year prior history with this topic**
- **Five “Hows” to Be Agile and Compliant**
- **Case Study of a Med. Device company’s Agile and Regulated Practices**
- **Your Questions**

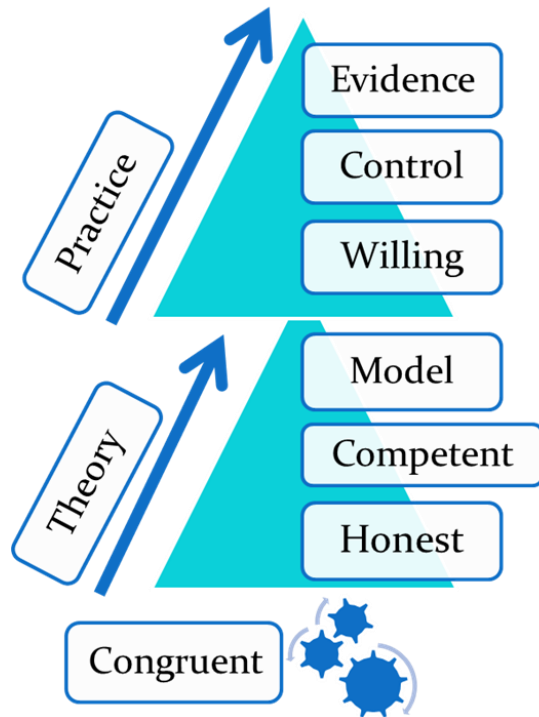
THE PROBLEM

*Conforming to Regulations while
Preserving Agile Practices*

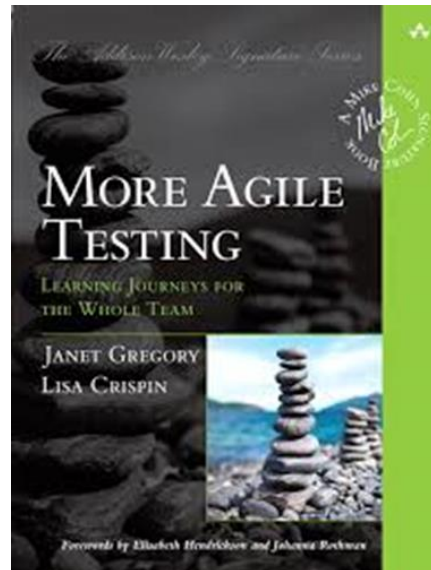
SOLUTION PATH

- Establish Specific Goals
- Define Deliberate Development
- Implement Risk Mitigation
- Generate Evidence and
- Close Gaps

SOME OF MY HISTORY



*What is Good
Evidence?*



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A FRAMEWORK

Theory

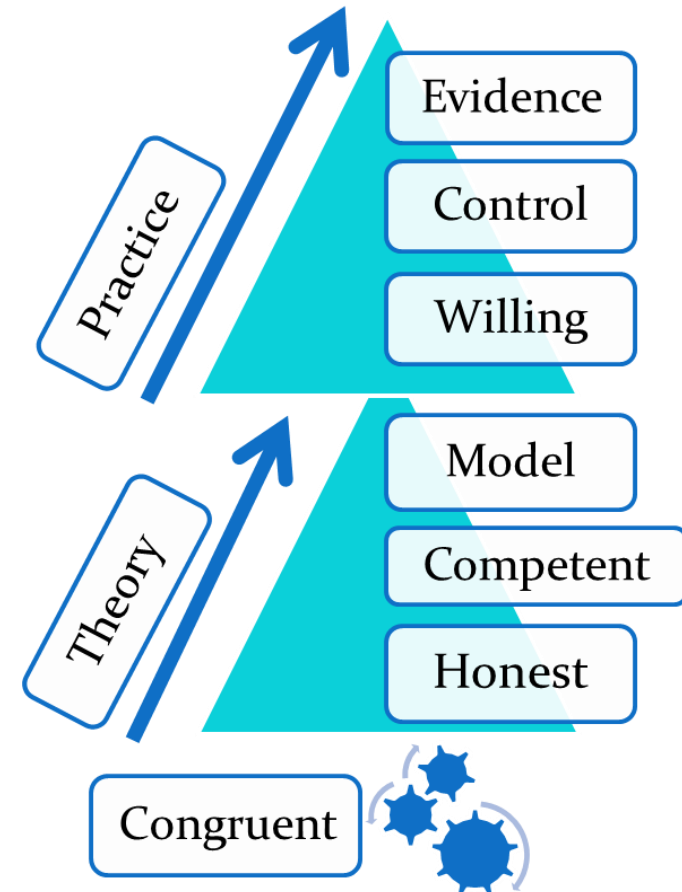
- Principles and Mindset

Practice

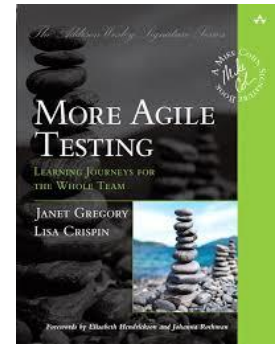
- Behaviors and Actions

...While Congruent

- Centered and Healthy



META-EXECUTION ...



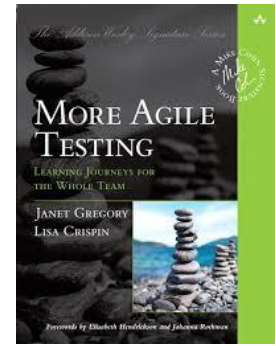
Regulatory Compliant Agile Testing

- Would stakeholders be happy if they saw/heard what I am doing right now?



- Is it important to share this information in the future?

... META-EXECUTION



Regulatory Compliant Agile Testing

- What is the most effective and efficient way to **Memorialize** this information – without breaking flow?
- Specific project “Hows” just become context specific details.



GOOD EVIDENCE

Regulatory Compliant Agile Testing

- Types and Qualities of Good Evidence

*What is Good
Evidence?*



May 2013

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THE FIVE HOWS ...

Establish Specific Goals

- How are you Regulated?
- What are the Expectations?
- Find your specific auditor's playbook



... THE FIVE HOWS ...

Define Deliberate Development

- Establish
 - Write, Draw, Explain - Development
 - Does it accomplish the Goal?
 - Are you capable of doing it?
- Maintain
 - Train and Sustain

... THE FIVE HOWS ...

Implement Risk Mitigation

- What can you foresee could go wrong?
 - Make a list, track, and manage it
- Be Reasonable and Prudent
- Nothing is Riskless
 - Some Risks gets accepted



... THE FIVE HOWS ...

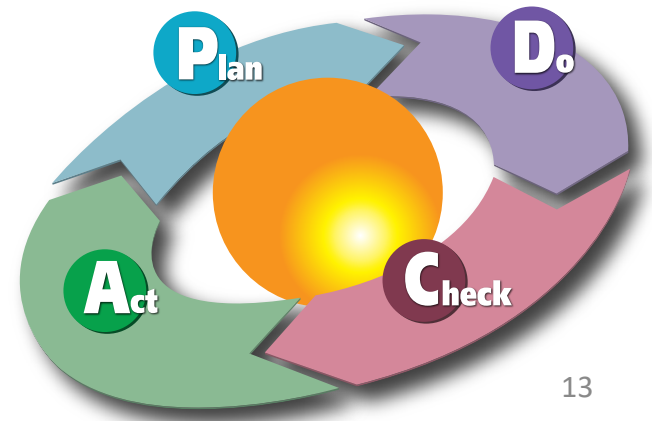
Generate Evidence

- If it is part of the official story
 - Memorialize it
 - Always add, never destroy
 - Include who, what, when, and why

... THE FIVE HOWS

Close Gaps

- Review and Revise
 - Retrospectives
 - Show Corrections and Improvements
- Show you are Under Control
- Don't require Perfect



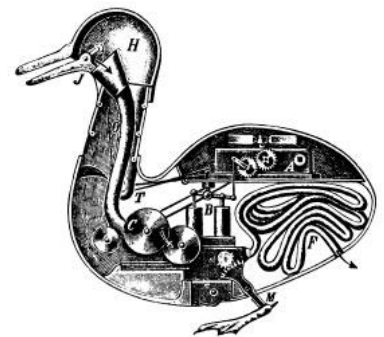
“MEDDEV” CASE STUDY



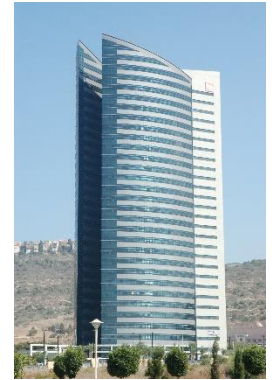
BEFORE WE START

System Thinking is needed

- Holistic approach focusing on the way that a system's constituent parts interrelate
- **Complex behaviors** over ever-smaller defined parts



MEDDEV



Company and Products

- Develops and manufactures **medical devices** used by doctors in hospitals
- Devices manage and change vast amounts of **diagnostic data**
- Several national and international **regulations**
- Internal project governance which **mandates** practices related to regulatory compliance via the **Quality System**

MEDDEV TEAMS

Teams

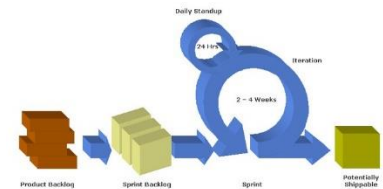
- Domain experts and software development
- About **130 team members** are organized into 8 scrum teams
- 50% employees, 50% are contractors
- Each scrum team is responsible for an **independent component** of MedDev's product



MEDDEV SCRUM

Scrum

- Using a scrum framework for **18 months** that inherited some practices from the former Waterfall approach
- Scrum team is **comprised of** programmers, testers, subject matter experts and analysts
- Each scrum team has a **scrum master** and a **product owner**



MEDDEV REG. COMPLIANCE

Regulatory Compliance

- MedDev has a team of regulatory compliance experts composed of a **dozen domain experts**
- Each expert has deep knowledge of relevant regulations for MedDev. Regulations may relate to product development, client onsite trials and system operations.
- RegComp experts act as **advisors** to Product Owners

MEDDEV REG. COMP. GROUP

Regulatory Compliance Group

- **Representative** to Agencies, Customers, and Suppliers
- RegComp establishes and maintains the MedDev **Quality System** and SOPs to the associated regulatory requirements and compliant practices
- RegComp **trains** everyone relating to regulatory compliance as expressed in the **Quality System**

MEDDEV QUALITY SYSTEM

Quality System

- The overarching controlling Policies and Procedures of MedDev
- Required by **Law**
- **Examined** by Governmental Agencies and Customers



MEDDEV SPRINTS

Sprints

- MedDev scrum sprints are **two weeks** long
- After each sprint **compliance evidence** is delivered to the RegComp
- This information is assembled into a package (**Design History File**) which is made available to internal and external auditors as needed
- **No special tasks** are required to create the regulated artifacts

MEDDEV DONE



Done Checklist

- Indicates what it means for a requirement to be completely implemented
- Constructed by the team, but **informed** by the Quality System and requirements from the RegComp group
- The product owner and the team agree on what it means for a story to be done

MEDDEV PRODUCT BACKLOG

Product Backlog

- The team product owner manages requirements with a product backlog
- Includes requirements which may be in the form of a user story
- May include a story that **encapsulates** a traditional specification
- Product Owner manages the product backlog

MEDDEV RELEASE PLANNING

Release Planning Event

- When a new project starts the scrum teams holds a release planning event
- During release planning, **regulatory risks** and regulatory practices are **reviewed**
- RegComp team is available to the scrum teams for **consultation**

MEDDEV BACKLOG REFINEMENT

Backlog Refinement

- Scrum teams have **standard** refinement sessions which may include RegComp to detail product backlog entries
- Refined backlog entries have may be tagged as needing **special** regulatory compliance attention, e.g., Link to Program Risks
- Story size is in Story Points

MEDDEV SPRINT PLANNING

Sprint Planning

- Standard negotiation re: what is in the Sprint
- Stories are decomposed into Tasks and Associated Hours
- Regulatory compliance is **built into** the normal work - capturing artifacts as required



MEDDEV DAILY STANDUP

Daily Standup

- Scrum teams have standard daily standup meetings
- Per their training, team members also **raise any concerns** they may have related to regulatory compliance
 - RegComp is then consulted

MEDDEV TRACEABILITY

Tracability

- Scrum teams **trace their software testing, programming activities to requirements and design artifacts**
- This information is **delivered** to the RegComp team at the end of the sprint

MEDDEV SPRINT DEMO

Sprint Demo

- Standard demo to stakeholders including members of the RegComp team



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MEDDEV RETROSPECTIVES

Retrospective

- Standard scrum team retrospective
- Suggestions that would modify or reinterpret the Quality System are a conversation with RegComp



MEDDEV RELEASE

Release

- Stories completed in the sprint require **no further software engineering work** and are potentially shippable product
- ...but there is a set of **additional 1 to 4 weeks** of Program Level activities performed by back office, non-scrum teams to transform potentially shippable product into **official shippable product**

MEDDEV VERSION CONTROL

Version Control and Archiving

- Per the Quality System, artifacts the scrum teams create and reference are in version control systems which can be **audited** (with zero notice) by members of the RegComp team and external stakeholders.
- The achieve is preserved for **17 years** after the last sale of the product

SUMMARY

Execute the Five “Hows” in your Context

- **Establish Specific Goals**
- **Define Deliberate Development**
- **Implement Risk Mitigation**
- **Generate Evidence and**
- **Close Gaps**

QUESTIONS AND STORIES



Delivering the Goods

***Harmonizing
Regulated and Agile Practices***



October 2018

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Thank you for attending this session.

Griffin Jones



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