INTEGRATING MEDICAL DEVICE PRODUCT DEVELOPMENT, DESIGN FOR SIX SIGMA AND QUALITY SYSTEMS REGULATION — PART I
By Dr Vinny Sastri

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Introduction
In today’s competitive, global environment, innovation and the development of new products are critical to a company’s growth and sustainability. New products that differentiate companies from their competitors serve as a competitive advantage and strategic strength.

Organic growth is the focus of many organizations that have spent the last few decades in intensive cost-cutting initiatives, processes and measures. A structured, flexible new product development process that facilitates speed to market, reduces product development cycle times, manages risks, and delivers products that meet the requirements of customers and the market place enhances the probability of product launch success.

For the medical device industry, documenting new product development activities from concept, design and manufacturing, to sales and distribution in a controlled manner is detailed in the Quality Systems Regulation (QSR) 21 CFR 820. Integrating the needs and requirements of the QSR into a corporate product development process can improve efficiencies, minimize risks and enable compliance to FDA and ISO regulations.

Part 1 of this article describes a new product development process that incorporates Design for Six Sigma (DfSS) tools. Part 2 will demonstrate how DfSS enables an organization to integrate the QSR requirements into the new product development process.

Best Practices for New Product Development
Dr Robert Cooper of MacMaster University, Canada, identified that only 11% of product development failures are for technical reasons associated with the performance of the product itself.1 The predominant causes of new product failure are marketing or market-related. These causes include the development of products the customer did not want and the development of me-too products. Cooper also identified that the single most important factor in determining a new product’s potential success was the presence of clearly superior product features that were well differentiated, offered significant customer benefits, and fulfilled unmet customer or market needs.

In April 2002, Industry Week published a survey on new product development. (See Table 1).2 In a March 2005 survey, Pete Collins from PriceWaterhouseCoopers found that 78% of fast growth CEOs mentioned new product development as a top priority that fit with strategic and corporate values.3 Developing new products has improved company revenues, earnings and profit margins, and has fundamentally changed their business processes.

About Winovia
Winovia® LLC, a consulting company that provides customized, sustainable solutions, strategies and training in new product development and quality management processes and high performance materials. Winovia employs the Six Sigma and Design for Six Sigma philosophy with the goal of strategic market penetration, improving product and process quality and increasing revenues and profits for its clients.

Winovia Specializes in:
- New Product Development Processes and Design for Six Sigma
- Manufacturing Processes, Process Validation and Six Sigma
- FDA and ISO Quality Management Systems for Medical Devices and Pharmaceuticals
- Strategic Technology Roadmapping

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Based on these studies, manufacturers can infer that success in new product development and innovation depends on several factors. A recipe for a successful product, therefore, should include the following ingredients:

- Clearly defined customer and product requirements up front, as well as customer collaboration.
- Management buy-in, support, and involvement with a clear business strategy and a shared vision.
- Product development considered as a business process, not relegated to technology and engineering alone.
- Dedicated resources and cross-functional teams (involving marketing, technology, production, sales, regulatory, legal, and purchasing) all the way from concept through commercialization.
- Clear metrics, relevant data, and data-driven decisions.
- A structured, flexible, quality, product development process from concept to commercialization for the entire organization.

### Table I. Key factors for successful new product development.

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<thead>
<tr>
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<th>Key Drivers</th>
<th>Time-to-Market Factors</th>
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<tbody>
<tr>
<td>Best fit with customer needs</td>
<td>69%</td>
<td>Collaboration with customers</td>
</tr>
<tr>
<td>Lowest product cost</td>
<td>13%</td>
<td>Allocation of resources</td>
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<tr>
<td>Innovative features</td>
<td>11%</td>
<td>Formal NPD process</td>
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<tr>
<td>First to market</td>
<td>5%</td>
<td>Shared platforms</td>
</tr>
<tr>
<td>Other</td>
<td>2%</td>
<td>Other</td>
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**Table I. Key factors for successful new product development.**

New Product Development Process

More than 70% of Fortune 500 companies have some sort of a formalized product development process. Insert figure with six different approaches to new product development. The goal of six sigma is to produce products with extremely low defects—3.4 defects in 1 million opportunities equivalent to a 99.9997% yield—based and trivialize the importance of vital but less-obvious activities. As a result, critical steps are missed, eliminated or overlooked. A new product development process is effective only if it includes all activities from the initial idea to final product sales.

The most commonly used process is the Stage Gate Process first introduced by Dr. Robert Cooper in 1988. It consists of five activity stages, between each of which is a decision-point or a gate for a go/no-go decision.

**Six Sigma**

Six Sigma is a business process that guides companies to produce high quality products, reduce costs, improve efficiencies and increase profits of existing products, processes and services. Six Sigma concepts can be included into an existing product development process or can be used to create a new product development process within an organization. Employing six sigma can also create profits of existing products, processes and services. First introduced by Motorola, the system relies on using rigorous data to drive decision making. By implementing the Six Sigma process, many companies like General Electric, Honeywell, Raytheon and IBM have realized billions of dollars in cost savings, process improvement and quality improvement.

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on those requirements that are critical to the customer. These requirements are called the critical-to-quality characteristics or CTQs. Figure 2 depicts a six sigma capable process and displays the relationship between defects, yields and the sigma level or sigma capability. Thus, a six-sigma product or process has 3.4 defects in one million opportunities, and a three-sigma product has 66,807 defects in one million opportunities, with a yield of 93.3%.

Defects can be quantified for any product, process, service or transaction. Rigorous root-cause analysis and statistical techniques can be used to identify the source of the defects, and corrective actions can be taken to implement a solution, and ensure that the defect never occurs again. The six sigma process is a five step process that is widely used around the world.

Define ▶ Measure ▶ Analyze ▶ Improve ▶ Control

Design for Six Sigma
The preamble to 21 CFR 820 specifies that manufacturers “…cannot inspect quality into the product, [they] have to build it into the design and process.”

By that token, device manufacturers can use DFSS as a tool for building quality into new products. DFSS is both a business management strategy and a product development process. It uses metrics, data and statistics, team dynamics, risk management, and project management tools to take products from concept to commercialization. The data-driven decision making process delivers a six-sigma capable product or service by focusing on design and process parameters based on customer and market requirements.

DFSS incorporates the Six Sigma data-driven principles into the new product development process. New products are designed, developed and distributed to six sigma quality and capability for those critical to quality requirements specified by the customer, market, or intended use.

DFSS uses a systematic approach. Manufacturers drill down from the customer’s requirements to detailed product properties and specifications. From there they consider product architecture and design (based on product requirements), and finally the detailed process controls that must be employed during production to ensure that a product meets the customer’s requirements (Figure 3).

These quantifiable drill-downs identify acceptance criteria, specifications and tolerances for the finished device or product and facilitate documentation to comply with the QSR.

Becton Dickinson (Franklin Lakes, NJ), a major medical device manufacturer, began using DFSS in its product development process in 2002. The implementation of DFSS by the company enables it to comply with the design control requirements of the QSR as part of its product development process.

The company underwent a significant cultural change to successfully adopt DFSS. In addition, training sessions and increased communications helped the process. The end result—a greater involvement with cross-functional teams, a clean line of site to end-use, and the ability to use the information for QSR compliance.

The DFSS process provides a series of tools that are especially suited to medical device manufacturing. Such tools include:

- **Portfolio management.** Organizations can assess product portfolios and formulate short-, mid- and long-term growth strategies.
- **Risk management.** DFSS identifies and addresses the risk at the end of each phase, enabling the teams and management to make informed decisions. Such practices can minimize potential risks and failures during production, sales and end-use.
- **Communication.** Standardized processes and templates facilitate communication between the team, the business and the customers. DFSS is very effective in working with remote and virtual teams.
• Project management. The structured product development process provides a clear understanding of the processes, deliverables, activities, timelines and budgets.

DfSS is not just about designing the product. The goal is that the new product, process or service should have robust (six sigma) performance in all stages. These principles should be seen by the customer, consumer, or end-user. They should be visible during product launch, as well as during production. In short, the six-sigma principles should be in the product by design. An example of a six-step new product development process is shown in Figure 4. There are six activity phases and six decision-making points or tollgates. The nomenclature in the figure is an extension of the six-sigma roadmap, which has five activity phases as follows:

- **Define** customer requirements, the problem, the project scope, and the customer goals.
- **Measure** the current product or process capability.
- **Analyze** what is wrong and identify possible root causes.
- **Improve**—find and test the possible solutions.
- **Control**—implement, monitor, and sustain the optimal solution.

The DfSS process was originally used for the design and engineering of complex machines and parts, where the overall product’s performance is dependent upon all of its sub-assemblies and components. The design specifications and capability of every individual component and sub-assembly is quantified, by a rigorous, step-by-step, drill-down from the finished product to every individual component. If each component and sub-assembly is six sigma capable within its specifications, it ensures six sigma capability of the final product (Figure 5). In the highly regulated medical device industry where performance and safety are critical, DfSS fits well for medical device product development. In the medical device scenario, each new product is designed so as to ensure defect-free performance—in other words, it is designed for it’s intended use with six sigma capability.

**Myths and Potential Barriers**

One of the biggest myths of DfSS is that it introduces extraordinary amounts of bureaucracy to the product development process. There is a perception that it increases timelines and delays product launches. In fact, the opposite is true. By identifying the characteristics that are critical to quality, DfSS focuses on the right scope, deliverables and the necessary activities of each stage providing speed and flexibility to the process. Teams are empowered to make decisions but they are also held accountable for those decisions. Decisions are made by assessing and addressing all potential risks.

Another misconception is that a structured product development process stifles creativity and innovation. DfSS actually stimulates innovation in product development by focusing on understanding the unmet needs of the market or the customer. Products that fulfill those unmet needs are the result of innovative solutions and execution.

Product development is all about bringing new products to market leveraging validated and proven technologies and core competencies. Long-term research that creates breakthrough and disruptive technologies should be conducted at a basic research level. These activities should be part of the long-term growth plan of the organization. Once the capabilities and potential of the breakthrough technologies have been validated, they can be used as platforms to spin off many new products for specific customers and applications using DfSS and new product development process.

The biggest hurdle for any initiative is the buy-in, support and involvement of upper management. There must be a shared vision from high-level managers. They must be integrally involved in the process and facilitate the cultural change. Results from such growth initiatives often take between two and three years to realize, compared to cost-cutting and productivity programs that realize benefits in six months to one year. The commitment must be sustained for the long foreseeable future.

**Benefits of Design for Six Sigma**

The benefits of a rigorous, data-driven product development process include:
Clearly defined product requirements based on intended use.

A fundamental understanding of the components and processes used to manufacture the device.

Differentiated and high-quality new products.

Cycle time reduction and speed to market.

Increased revenues and profits.

Flexibility to respond to new requirements, applications and regulations.

Better assessment of risks through the product development cycle.

Sustainable products and processes.

**Conclusion:**

DfSS can be integrated into a company’s existing product development process. It uses a systematic, data-driven approach to quantify critical product requirements based on the intended use. Products are designed and developed by fundamentally understanding how components, sub-assemblies and processing quantifiably relate to the performance of the finished device.

Finished devices can be designed, developed, manufactured, and sold with six sigma quality and capability meeting the end-user needs for safe and effective products. Implementing the concept, leads to product innovation, consistency, and reliability, as well as sustainable sales, revenues and profits.

**REFERENCES**


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**Is Design for Six Sigma Right for You?**

In order to assess whether or not Design for Six Sigma is right for you or your organization here are some questions you should want to answer. (Use a scoring system of 1 – 10: 1=very poor, 5=fair, 10=superior, or 1=never, 5=sometimes, 10=always)

1. How good is your organization’s product development process?

2. How often does the scope of the product requirements change?

3. How well do you assess customer needs and intended use?

4. How well do you use quantifiable metrics for the new product specifications that relates quantifiably to the intended use?

5. How well do you drill-down to key component characteristics and specifications with a line-of-sight to the intended use?

6. How often do you use predictable tools to assess your product capability by design?

7. How well do you understand the raw material or component specifications and processing controls as it relates to the product performance and the intended use?

8. How often do you use product capability scorecards to measure product performance?

9. How good is your risk management system across the product development cycle?

10. How good are your product launches (the right product, with the right requirements, no tweaking and refining after launch, etc.)?

Any score below 80 suggests that you should reevaluate your product development process and include more rigor within it.