

14-Sep-2015

Before defining the quality it is better to have a look on the dimensions of quality.

Dimensions of Quality:-

Garvin 1987 provides an excellent discussion of 8 components or dimensions of quality. Following are the components.

- 1.- Performance
- 2.- Reliability
- 3.- Durability
- 4.- Service-ability
- 5.- Aesthetics
- 6.- Features
- 7.- Perceived quality
- 8.- Conformance to Standards.

1. Performance:-

Will the product do the intended job.

2. Reliability:-

How often does the product fail.

3. Durability:-

How long does the product last.

4. Serviceability:-

How easy is it to repair the product.

5. Aesthetics:-

What does the product look like.

6. Features:-

What does the product do.

7. Perceived Quality.

What is the reputation of company or its products.

8. Conformans to Standard.

Is the product made exactly as the designer intended.

Develop a strategy for launching a new brand in your locality or area. 15-Sep-2015

On university road I want to prefer a to open an ice-cream shop where self service is provided. These shop have different flavour of ice-cream & customer make his himself as he desire.

On in university there may be a bakery "Cakes & Bakes" open.

29. Sep. 2015

Definitions of Quality:

From the discussion it is clear that quality is indeed a multi-facet entity. Consequently a simple answer to a question What is quality? or what is quality improvement is not easy.

The traditional definition of quality is based on view point that:

The products & services must need the requirements of ^{✓ who} those to use them. i.e.

"Quality is fitness for use"

Generally there are two aspects of fitness for use

- (i) Quality of design
- (ii) Quality of conformance

(i) Quality of design:

All goods & services are produced in various grades or levels of quality these variations in grades are ~~ent~~ intentional & consequently the appropriate technical term is quality of design. e.g.: All auto-mobils differ with size appearance, performance etc.

(ii) Quality of Conformance:

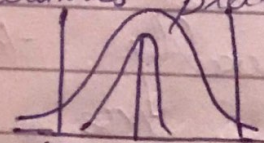
It is how well the product conforms to the specifications required by the design. Quality of conformance is influenced by a number of factors e.g.: training & supervision of the work force, test & inspection activities etc.

The modern definition is so we will this definition:

"Quality is inversely proportional to its variability"

$$\sigma \propto \frac{1}{V}$$

e.g. Japan's products & other countries products



13-Oct-2015

Quality engineering terminology:

Every product possess a number of elements that jointly described what the user or consumer thinks of as quality. These parameters are often called quality characteristics. Quality characteristics may be of several types:

- Physical
- Sensory
- Time orientation

1. Physical:

Length, weight, voltage. Things that can be measurable or touchable.

2. Sensory:

Taste, appearance, colour etc

3. Time Orientation:

Things that can be measurable w.r.t time. e.g Reliability, durability, serviceability.

Note:-

Note that the different types of quality characteristic can relate directly or indirectly to the dimensions of quality.

Most organization find it difficult to provide the customer with products that have quality characteristics that are always identical from unit to unit or are at levels that match customer expectations.

A major reason for this is variability. There is a certain amount of variability in every product & consequently no two units of products are ever identical.

If this variation is small then it may have no impact on the consumer. However, if the variation is large then the customer may perceive the unit to be undistressable & unexpected.

Sources:

Sources of variability include differences in material, differences in performance & operation of the manufacturing equipment & differences in the way the operators perform their task.

Statistical Methods:

The variability can only be described in statistical terms. Statistical methods play a central role in quality improvement efforts. To apply statistical methods it is fairly typical to classify data on quality characteristic as either attributes or variable data.

Variable data:

Variable data are usually continuous measurement such as: length, speed, life, voltage etc.

Attribute data:

Attributes data on the other hand are usually discrete data often taking the form of counts such as: no. of defective fans.

Quality characteristics are often evaluated relative to specifications.

Specifications:-

Specifications are desired measurement for the Q.C of the components that make up the product as well as the desired values for the Q.C in the final product

Target:-

A value of a measurement that corresponds to the desired value for Q.C is called nominal or target value for that characteristic.

USL:-

The largest allowable value for a Q.C is called upper specification limit. USL

LSL:-

The smallest allowable value for a Q.C is called a lower specification limit LSL

⇒ The specification limit control by engineers.

→ Read usel) conforming, non-conforming, defective, non-defective, conformity & conformities, defect & defects.

Non-conforming products:- "are those that fail to meet one or more of its specifications."

Non-conformity:- "A specific type of failure is non-conformity" e.g:- kam ho raha ha.

⇒ A non-conforming product is considered defective if it has one or more defects.

- ⇒ What is Statistical process control?
- ⇒ Chance causes of variation & assignable causes of variation
- ⇒ Statistical basis of a control chart.

Statistical process Control:-

Process:-

Process is a sequence of inter-dependent & linked procedures which at every stage consume one or more resources (employees, time, energy, machines, money etc.). to convert inputs (materials, different parts etc.) into outputs

OR

Process is a series of actions or steps taken in order to achieve a particular role or purpose

OR

Process is a mechanism to generate outputs by giving inputs. eg:- car mechanism, comp u
input something to get output

Note:-

Mostly processes are repeatable

Control:-

Control is a way to identify the unusual changes in the process & then elimination of these changes & hence bring the process into normal state.

SPC:-

It is a powerful collection of problem solving tools useful in achieving stability & improving capability through the reduction of variability.

SPC is one of the greatest technological developments of the 20th century following are the major tools of SPC.

- 1- Histogram OR Stem & leaf Plot
- 2- Check Sheet
- 3- Pareto chart (80-20 distribution e.g. in economic view 20% people control 80% things)
- 4- Cause & effect diagram
- 5- Defect concentration diagram
- 6- Scatter diagram (see the direction of two variables)
- 7- Control chart

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Chance causes & Assignable causes of variations:-

In any production process regardless of how well designed & carefully maintained it is a certain amount of inherent or natural or variability will always exist. This natural variability or background noise is the cumulative effect of many small essentially unavoidable causes. In statistical quality control this natural variability is called a stable system of chance causes.

A process that is operating with only chance causes of ^{variation} operation is said to be statistically in-control. In other words

chance causes are an inherent part of the process.

Other kinds of variability may occasionally be present in the output of process. This variability in quality characteristic usually arises from the following sources

- (i) Improperly adjusted or controlled machines
- (ii) Operator error or defective raw materials

Such variability is generally large when compared to the back-ground noise & it usually represents an ^{above} ~~un-~~expectable level of process performance. We refer to these sources of variability that are not part of the chance causes as assignable causes of variations. A process that is operating in the presence of assignable causes is said to be an out of control process. A major objective of SPC is to quickly detect the occurrence of assignable causes of process shifts. So that investigation of the process & corrective action may be undertaken before many non-conforming units are manufactured.

To monitor location parameter μ & dispersion parameter σ . Walter A. Shewhart introduced the concept of control chart.

Chance causes / Common causes of variation / in-control
 Assignable causes / Special causes of variation / out of control

General Model for Shewhart Model chart: 26-Oct-20

Let w be a sample statistic that measure some quality characteristic of interest & suppose that mean of w is " μ_w " i.e. $E(w) = \mu_w$ & the standard deviation of w is σ_w i.e.

$$S.D(w) = \sigma_w = \sqrt{E(w - E(w))^2}$$

Then the central line (CL), the upper control limit (UCL) & the lower control limit (LCL) become

$$UCL = E(w) + k S_d(w) = \mu_w + k \sigma_w$$

$$CL = E(w) = \mu_w$$

$$LCL = E(w) - k S_d(w) = \mu_w - k \sigma_w$$

Where k is the distance of the control limits from the central line expressed in standard deviation units. This general theory of control charts was first proposed by Walter-A. Shewhart & the control charts developed according to these principles are often called Shewhart control charts.

The most important use of a control chart is to improve the process. Following observation generally have been made.

- (i) Most processes don't operate in a state of statistical control & consequently the you'll see & attentive use of control chart will identify assignable causes if these causes can be eliminated from the process, variability will be reduce & the process will be improved.
- (ii) The control chart will only detect assignable causes.
- Management, Operator & Engineering action will usually be necessary to eliminate the assignable causes.

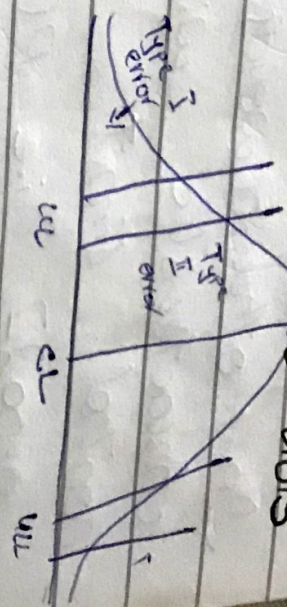
Control Charts

= $n \sin(\theta)$ (rand)

27-Oct-2015

Process. Control or indicate some

3rd - Nov-2015



If we widen the limits then Type II error increases & Type I error decreases

Choice of Control Limits:-

The critical decision that must be made in designing a control chart. By moving the control limits further from the central line we decrease the risk of a type I error that is the risk of a point following

Falling beyond the control limits indicating an out of control condition when no assignable cause is present.

However, widening the control limits will also increase the risk of Type II error that is the risk of a point falling b/w the control limits when process is really out of control. If we move the control limits close to the central line the opposite effect is obtained.

Consider the quality characteristic say X follows normal distⁿ with mean μ & variance σ^2 then the following properties holds if we fix $k=3$

$$P(\mu - 3\sigma < X < \mu + 3\sigma) = 0.9973$$

In other words we say that the quality characteristic will fall beyond the control limit. Fixing $k=3$ we have the prob.

$$0.0027 \text{ i.e. } P(X > \mu + 3\sigma \text{ or } X < \mu - 3\sigma) \\ = P(X < \mu - 3\sigma) + P(X > \mu + 3\sigma) = 0.0027 \rightarrow \text{level of significance}$$

This prob of 0.0027 is known as prob. of type I error or false alarm rate (FAR). Prob of type I error is also known as FAR in quality control perspective.

In other words

$$P(X > \mu + 3\sigma) = 0.00135$$

$$P(X < \mu - 3\sigma) = 0.00135$$