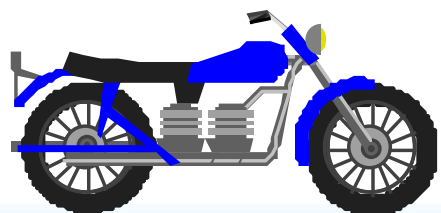
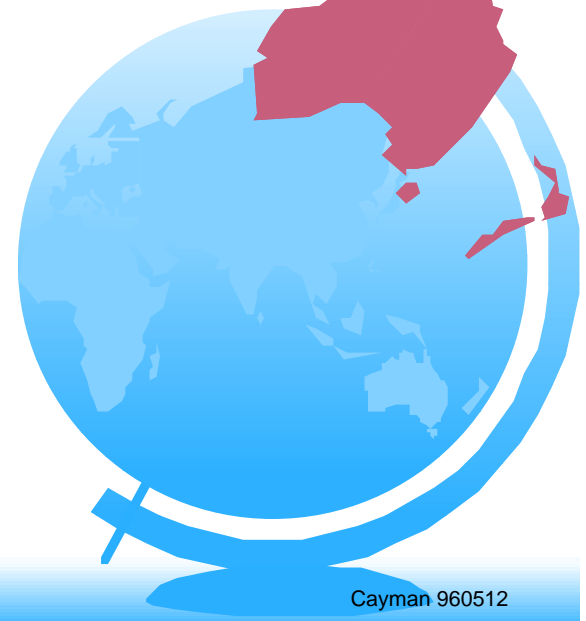




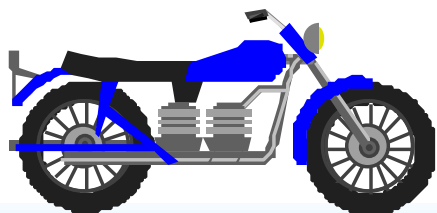
# ISO 9000

## An Overview



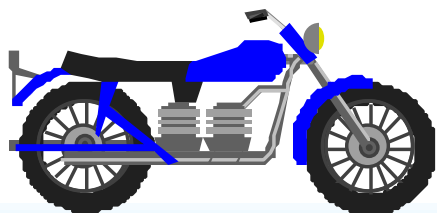
# Introduction

- ❖ The Purpose of this Presentation is to Provide an **Overview** of **ISO 9000** and **What It Means to Everyone at Company X**



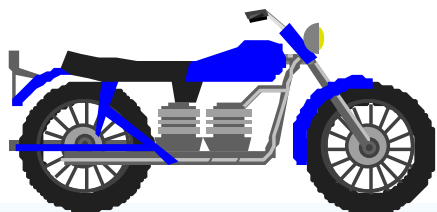
# ISO 9000

- ❖ The Ultimate **Goal** of ISO 9000 is to Provide **CONSISTENT PROCESSES**
- ❖ Documented Systems Provide For Consistency
- ❖ **RESPONSIBILITIES** Defined
- ❖ Periodic **Internal & External Audits** Ensure Systems Are Working



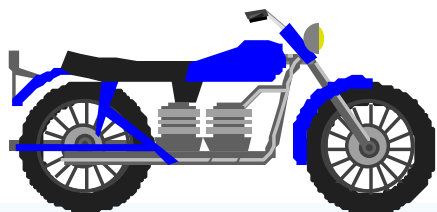
# ISO 9000 Origin & Scope Of Application

- ❖ International Organization for Standardization
- ❖ TC 176 - Meets in Geneva, Switzerland
  - TC = Technical Committee
- ❖ Europe Wide
  - Must Be Registered to Sell In Europe
- ❖ Japan is Accepting
- ❖ US Military Switching From MIL-Q-9858A



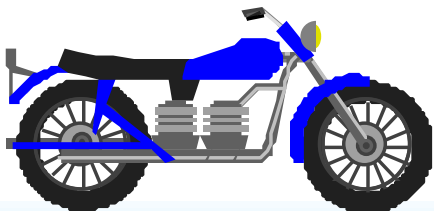
# ISO 9000 Documents

- ❖ ISO 9000-1
  - Quality Management and Quality Assurance Standards - **Guidelines for Selection and Use**
- ❖ ISO 9001
  - Quality Systems - Model for Quality Assurance in **Design, Development**, Production, Installation and Servicing
- ❖ ISO 9002
  - Quality Systems - Model for Quality Assurance in **Production, Installation and Servicing**
- ❖ ISO 9003
  - Quality Systems - Model for Quality Assurance in **Final Inspection and Test**
- ❖ ISO 9004
  - Quality Management and Quality System Elements - **Guidelines**



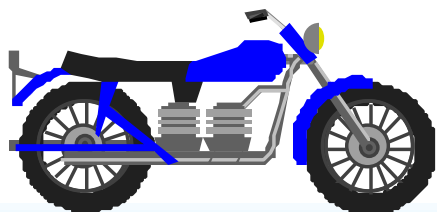
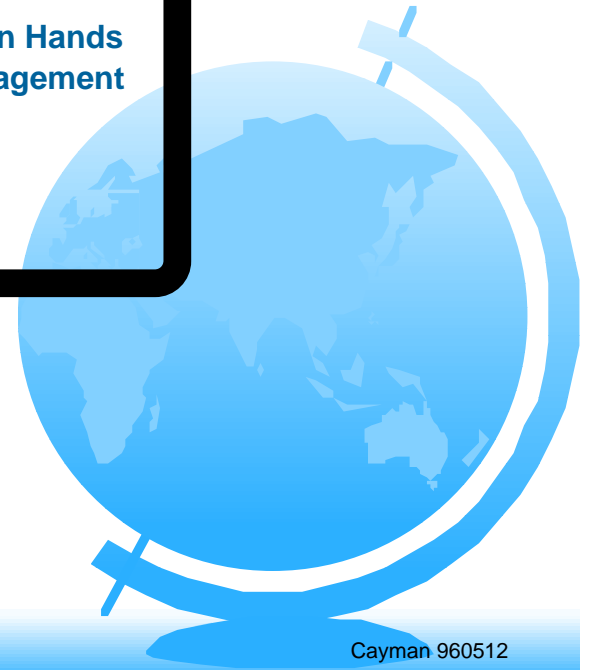
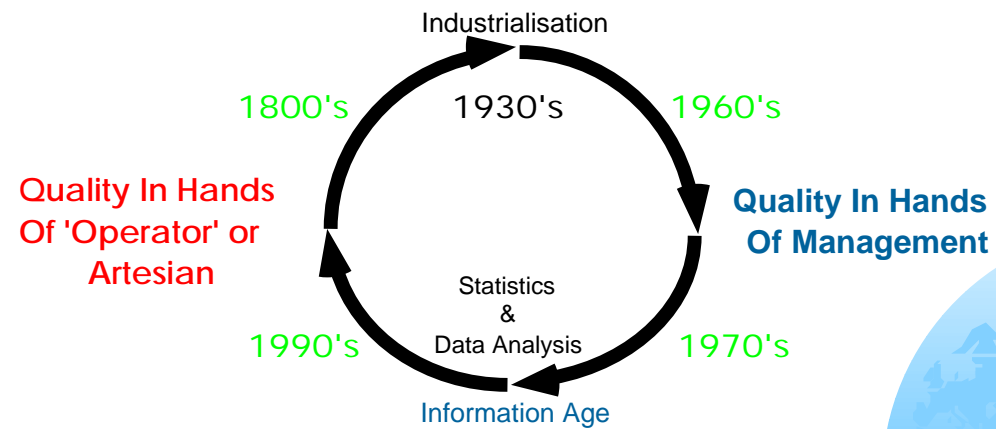
# ISO 9001 Sections

- ◆ 4.1 Management Responsibility
- ◆ 4.2 Quality System
- ◆ 4.3 Contract Review
- ◆ 4.4 Design Control
- ◆ 4.5 Document & Data Control
- ◆ 4.6 Purchasing
- ◆ 4.7 Control of Customer Supplier Product
- ◆ 4.8 Product Identification & Traceability
- ◆ 4.9 Process Control
- ◆ 4.10 Inspection and Testing
- ◆ 4.11 Control of Inspection, Measuring and Test Equipment
- ◆ 4.12 Inspection and Test Status
- ◆ 4.13 Control of Non-Conforming Product
- ◆ 4.14 Corrective and Preventative Action
- ◆ 4.15 Handling, Storage, Packaging, Preservation & Delivery
- ◆ 4.16 Control of Quality Records
- ◆ 4.17 Internal Quality Audits
- ◆ 4.18 Training
- ◆ 4.19 Servicing
- ◆ 4.20 Statistical Techniques



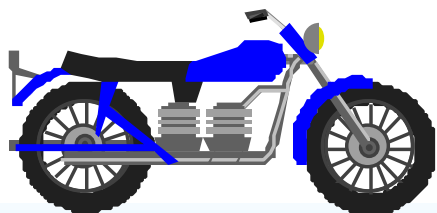
# Responsibility

## Circle (or Cycle) Of Responsibility



# Documentation

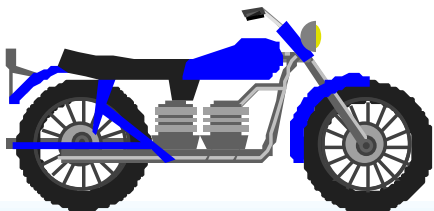
- ◆ Organisation Charts
- ◆ Procedures
  - Flow Charts
  - Process Sheets
  - Process Routers
  - Work Instructions
- ◆ Forms
- ◆ Tags
- ◆ Prints
- ◆ Specifications
- ◆ GES Specs
- ◆ SPC Data
- ◆ Inspection & Test Results





# Myths vs. Truths

- ❖ This Is **NOT** An Effort To **CHANGE** The Way You Do Things Now
- ❖ Documentation Is Meant To Be **Easily Changed**
- ❖ The **Less** Documentation, The **Better**



# Basic Rules

## ❖ Your Job & Documentation

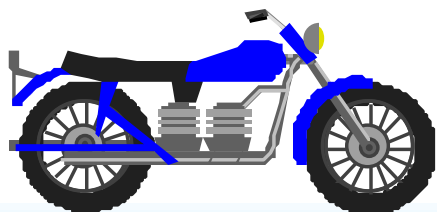
- **SAY** What You **Do**

  - ◆ Documentation

- **DO** What You **Say** You Do

  - ◆ Actions

## ❖ If It's **Not WRITTEN** Down, It **DIDN'T** Happen

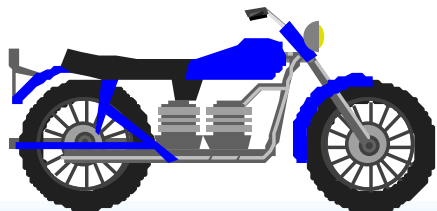


# Real Life

## What ISO 9000 Means To You

### ❖ You **MUST**:

- Know Your **Job Duties**
- Know What **Training Your Job Requires**
- Be Able To Tell **About How You Were Trained**
- Know **What Documentation Involves YOU!**
- Know **How To Find Out** What The 'Latest' Version Is
- Know **What The Documentation Says**
- Know **How** The Documentation **Applies To YOU!**
- Know What The **INTENT** of the Documentation

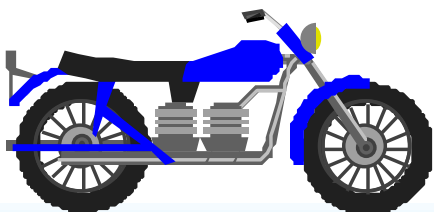


# Quality Policy

❖ You MUST Know **What** The **Company**  
**X Quality Policy Is**

And

❖ You WILL Be Asked **What** The **Quality**  
**Policy Means To YOU!**



# Things To Be Alert For

- ❖ Container Labelling
- ❖ Documents - Watch for “Headers”
  - Title
  - Date
  - Signature or Initials
  - Originating Department or Group
- ❖ Calibration Labels
  - Measurement and Test Equipment
  - Assembly Equipment (e.g.: Torquing Tools)

