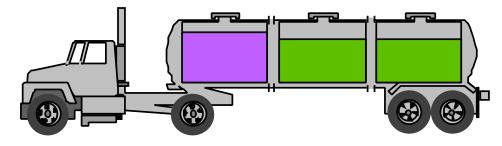


# ISO 9000 In The Over-The-Road Transportation Industry



#### Schedule

8:45 AM - Part 1

9:30 AM - Break

5 Minutes

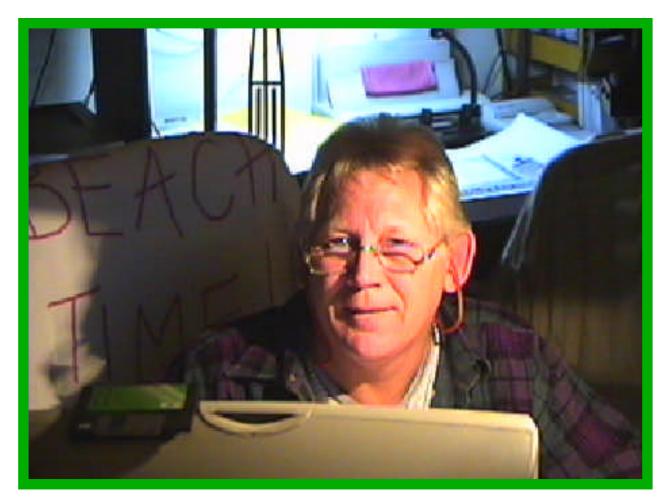
9:35 AM - Part 2

10:30 AM - Break

15 Minutes

10:45 AM - Part 3

11:30 AM - End Of Session



# Hello! I'm Marc and I'm glad you're here!

#### **Topics**

- ISO 9000 Generalities
- Who's doing ISO and Why
- About Auditors
- A brief 'How to get started' guide
- ISO 9000 Some Specifics
- ISO 9000 Element by Element
- Cautions and Wrap-Up

#### The Basics

#### ISO Boiled Down To Its Base

#### Say What You Do

This means document your systems so you will consistently do the job the same way every time. We must make sure we have appropriate documentation.

#### Do What You Say

This is what the auditors want to see. Objective evidence that what you say you are doing in your documentation is what you are doing in practice.

#### Technically, What Is ISO 9000 About?

- Quality Management
- Quality Assurance
- Quality System Deployment and Documentation
- Records and Information Management Play an Integral Role
- Has Little to do with Quality
- Does NOT Demonstrate Quality of Product
- Should Result in a Better Product and/or Service

#### What About QS 9000?

- Transportation companies are not subject to QS 9000 registration.
- QS 9000 is a requirement of Ford, GM and Chrysler.
- The 'Big Three' are requiring most carriers to obtain ISO 9000 registration.
- Each has it's own specific supplier requirements.
- You \*may\* be subject to QSA-S "Quality System Assessment for Services" (Ford).

#### Federal Rules & Regulations

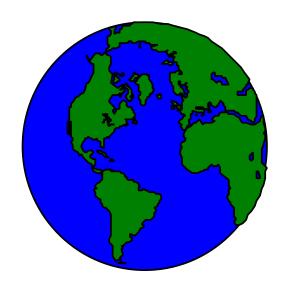
- ISO 9000 does not change or affect any federal rules or regulations.
- Federal requirements are \*in addition\* to ISO requirements
- Systems documentation will weave together all requirements

# Global Competition Causes Surging 9001 and 9002 Registrations

- Approximately 50% of all US firms will be registered by the end of 1998 (WSJ)
- Europe wide EU (EEC)
- Pacific Rim embracing

#### Who's Doing ISO?

- European Economic Community
- The 'Big Three'
- DoD
- NASA
- Over 80 countries have embraced the ISO Series of standards as National Standard





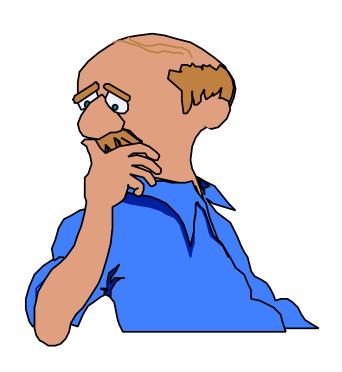
#### Why ISO 9000 Registration?

- Improves Internal Operations
- Enhances Competitive Position
- Required for International Business
- Customer Satisfaction
- Foundation for a Quality Program
- Complement Existing Programs
- Market Share
- Competitive Pressures



#### Why Do Companies Register?

- Customer
   Requirement (>80%)
- Sales and Marketing 'Tool'
- NOT for Quality
   Improvement



#### Internal Effects

Effective, comprehensive quality management system

Blueprint for building quality into products and services

Increase potential growth, competitiveness and profit

Consistency of product and services leads to happier, more effective employees and pleased customers

#### Critical Success Factors

- Dedicated 'Company Knowledgebase'
   (Co-ordinator and/or Management Representative)
- Pre-assessment
- Involved, supportive top management
- Receptive culture
- Focus on business rather than functional areas
- Prioritise processes based on customer needs, anticipated benefits, and potential for success

#### Registration

- Your company will choose a registrar.
- Registration lasts for 3 years.
- Registration is NOT a 'one-shot' deal.
- Your company will be re-audited at least once a year.

#### Forever.

And ever.

And ever..... (Not a fairy tale!)

#### **Auditors!**

 Registration requires regular audits by your Registrar. These are called Third Party audits. Just as has been done in banks for years, auditing has reached every industry. Whether twice a year or once a year, your company quality system has to audited by the company which registers your company. That company is your Registrar.

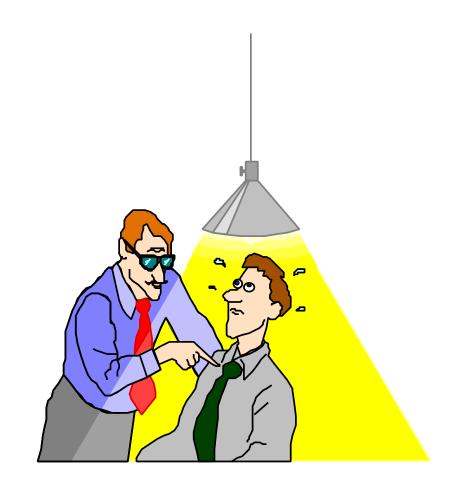


#### **Reasons For Audits**

- ISO 9000 Requires Them (4.17)
- A Control Mechanism Used By Management
- Tool For Continuous Improvement
- Correct Nonconformities In Systems
- Helps Ensure Ongoing Systems
   Operate As Intended And As Required

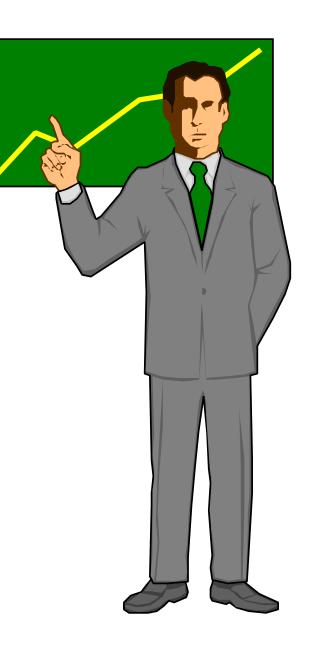
#### Auditors Are Not!!!

- Inquisitors
- Fault Finders
- Rock Throwers
- Avenging Angels (Biased For or Against)
- Dishonest
- Overactive



#### Payback

- Companies minimize deficiencies in supply and support of products and services.
- Companies identify problem areas and address them quicker.
- Companies identify customer needs more accurately.
- Companies become more consistent in their product and services.



# OK, Already. Which Way Do We Go From Here? I thought this was a 'How To....' Seminar!

#### First, Some Considerations....

- Your Product is Transportation of 'goods'
- Your Product is (technically) a service
- Customer Supplied Product may include a customer's trailer
- Your SIC Code is

4200 - Motor Freight Transportation & Warehousing

Sub-categories include

4212 Food "Containers Frozen Food Transportation"

4212 Live 1 "Livestock Transportation"

4213 Tran 2 "Transportation and Distribution Management"

#### General Registration Path

- Assess your situation (Pre-assessment)
  - May be by registrar or consultant Also called Gap Analysis
- Define a plan with time line & begin
- Interview and choose registrar
- Documentation processes
- Manage transitional activities
- Registrar document review
- Registrar pre-assessment
- Corrective actions
- Registration audit
- Implementation timeframe: 3 to 9 months

#### Typical Registration Costs

- Office Staff = 15 to 30 Souls
- Costs spread over 3 years

•	Registrar	US\$10K

<ul> <li>In-House Resources</li> </ul>	US\$15K
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<ul> <li>Consultant</li> </ul>	US\$15K
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**US\$45K** 

Note: Range = US\$15K to US\$100K (\*Plus)

#### Failure Modes

- Lack of documentation
- No or inadequate document control
- Poor record keeping and systems
- Lack of management involvement
- Poor communication
- Personnel not following documentation

#### A Word About Registrars....

- Choose several registrars and interview them each in-house
- Determine their man-day fees and hours (some cite 6 hours as a 'man-day')
- What is included in their 'expenses'?
- Negotiate/bargain for your best deal
- Determine their 'Scope of Authority'
- Are their auditors on contract or are they hired as Full-Time employees?

# Now - Practically Speaking, How Do We Start?

#### Appoint a Champion

This person will be your company 'Knowledge Base'. Begin 'Document Identification and Mapping'

#### Where are you at now?

The Champion should do a self analysis of the company, begin to develop an implementation plan and should begin to contact registrars.

#### Call a Meeting!!!

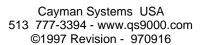
Do a roll-out! Bring everyone together and explain the basics of ISO and what it means to each person and to the business. Give 'Marching Orders'!

#### Make a Project Plan!!!!

					1996	1997
Task ID	Project Elements	Days	Start	Finish	Feb Mar Apr May Jun Jul Aug Sep Oct Nov Dec	Jan Feb Mar Apr May Jun Jul
1	Determine Specific Requirement(s)	1d	2/1/96	2/1/96		
2	Define Time to Complete Requirement	1d	2/1/96	2/1/96		
3	Define Scope of Assessment	1d	2/1/96	2/1/96		
4	Project Set-Ups	178d	2/1/96	10/7/96	▼	
5	Write Company Quality Systems Manual	1d	2/1/96	2/1/96		
6	Document Company Quality Policy	1d	2/1/96	2/1/96		
7	Define Documentation Systems	1 d	2/1/96	2/1/96		
8	Document Master Numbering System	1d	2/1/96	2/1/96		
9	Establish Master Binders	178d	2/1/96	10/7/96	▼	
10	Procedures History Binder	1d	2/1/96	2/1/96		
11	Project Master Binder	1d	2/1/96	2/1/96		
12		1d	10/7/96	10/7/96	0%	
13	Contact Registrar	157d	3/1/96	10/7/96	<b>▼</b>	
14	Agree on Scope	1d	3/1/96	3/1/96	% <b> </b>	
15	Agree On Fees (Try to Bargain)	1d	3/1/96	3/1/96	% <b> </b>	
16	Agree On Audit Date(s)	1d	3/1/96	3/1/96	% <b> </b>	
17	Submit Required Documentation	1d	3/1/96	3/1/96	% <b>I</b>	
18		1d	10/7/96	10/7/96	0%	
19	Awareness & Information Meetings - Hourly	198d	2/1/96	11/4/96	▼	
20	ISO 9000 Awareness	1d	2/1/96	2/1/96		
21	Work Instructions & Documentation	1d	5/3/96	5/3/96	0%	
22	Auditee Training	1d	8/5/96	8/5/96	0%	

#### How to Start - Part 2

- The Champion should start flow charting and understanding the 20 master systems and a quality manual should be \*started\*
- Individuals should begin listing and flow charting their jobs
- Do you need a Piano Teacher?
- This is a step by step building process
- The Champion orchestrates



#### A Word to the Wise...

- Your quality systems manual is one of your most important documents
- 'Canned' texts will have to be tailored
- Best way is to take the ISO 9000 text and tailor it as you develop your internal systems for each ISO element
- DO NOT try to write your quality manual and then write procedures to support it

#### How to Start - Part 3

- Do another self assessment when you have everyone's input
- If you think you're ready, arrange for the registrar's document review
- Next will come pre-assessment by the registrar
- Registration audit
- \*\* It takes 2 to 3 months to get your certificate after your registration audit

### ISO 9000

## The Specifics

#### Foundation of ISO 9000 Series

- The linking thread through out the ISO 9000 standards is the emphasis on recording information that pertains to all aspects of quality and management.
- While there are several explanations of the reason for the origin of the series, the most basic reason for their coming into existence relate to ensuring responsibilities are defined for Liability Issues.

#### General Terminology

Registrar

Must be accredited (RAB, minimum)

Registration Audit

An auditor comes and reviews documentation and practices

- Companies obtain registration and receive certificates
- Auditor

Must be certified by the RAB (minimum)

#### **ISO In Detail**

- International Organisation for Standardization
- isos from the Greek word meaning equal
- Founded 1946
- Based in Geneva, Switzerland
- ISO 9000 series begun about 1978
- Developed initially to support two-party contractual agreements (has expanded)
- ISO 9000 series released in 1987
- Over 80 countries have embraced as a National standard

#### ISO In Detail II

- International Standards on Quality
   Management and Quality Assurance
- Established by ISO Technical Committee 176 (TC176)
- Used by both manufacturing and service industries
- Recognised World-Wide:
  - International Seal of Approval
- Defines an effective quality system

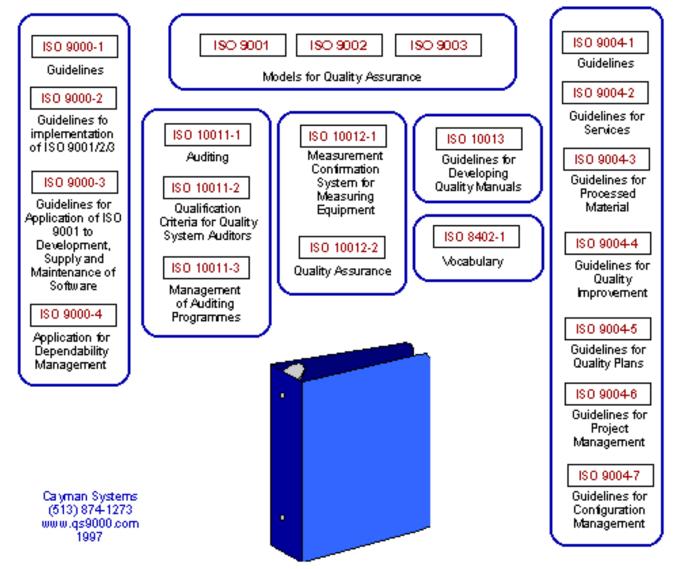
#### ISO In Detail III

- Requires documentation for quality system elements
- Requires 3rd party audits and registration
- Applies to all kinds of manufacturing and all kinds of service organizations
- Establishes consistent quality system practices that cross international borders
- Reduces or eliminates customer audits
- Provides a common language and set of terms

## Scope of the Standards

- Clarify quality-related principles and concepts
- Provide guidelines
- Specific requirements to achieve customer satisfaction by preventing nonconformances
- Inspection and test conducted on finished products and services can be satisfactorily demonstrated

### ISO 9000 Documents



Management and Quality Assurance - Vocabulary

This document is an attempt to provide consistent definitions for major terms.

This document is important to the interpretation of the ISO 9000 Series of documents.

Quality Systems - Model for Quality Assurance in Design, Development Production, Installation and Servicing

This is the main ISO 9000 Series document. It's contents guide the entire series. It is typically the document businesses register to. It contains provision for design development and control.

Quality Systems - Model for Quality Assurance in Production, Installation and Servicing

ISO 9002 is the exact same document as ISO 9001 EXCEPT that design is not included. This is the model transportation companies register to.

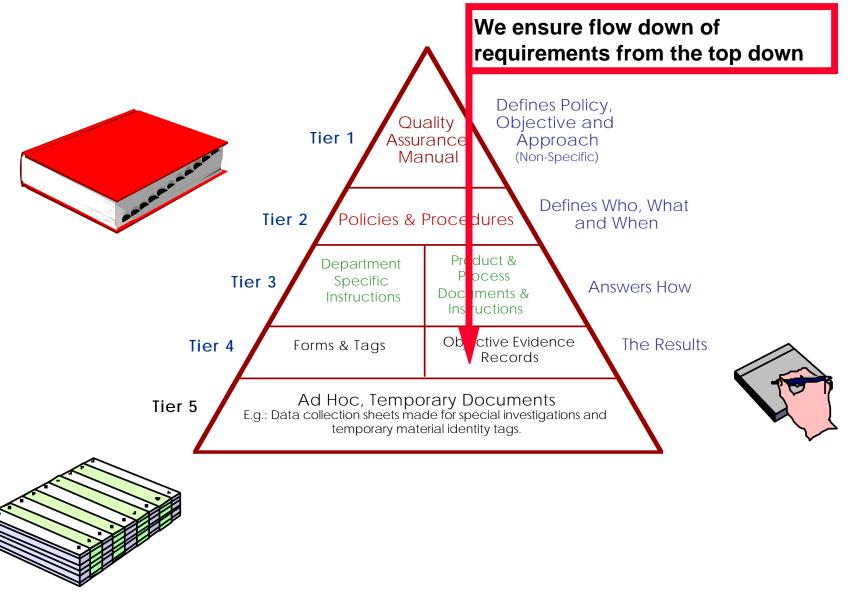
Quality Systems - Model for Quality Assurance in Final Inspection and Test

ISO 9003 is essentially a dead document. It's intended sector focus has embraced ISO 9002 or ISO 9001.

## Quality System Requirements

- Quality Policy
- Quality Manual
- Procedures
- Work Instructions
- Other Documentation
- Internal Auditing

### ISO 9000 Document Structure



### The Elements of ISO 9001

#### Each document is divided into 4 sections

- 1. Scope
- 2. Normative Reference
- 3. Definitions
- 4. Quality System Requirements

Section 4 contains the requirements which is why you will hear a lot of people cite the standard by section and sub-section, such as "4.3 Contract Review"

### List of Elements

4.1	Management Responsibility
4.2	Quality System
4.3	Contract Review
4.4	Design Control
4.5	Document and Data Control
4.6	Purchasing
4.7	Control of Customer-Supplied Product
4.8	Product Identification and 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 Preventive Action
4.15	Handling Storage Packaging Preservation and Delivery
4.16	Control of Quality Records
4.17	Internal Quality Audits
4.18	Training
4.19	Servicing
4.20	Statistical Techniques

## 4.1 Management Responsibility

- Management Responsibility is applicable to every company registering to ISO 900x. It can be one of the most difficult to achieve when management fails to support and involve themselves in the effort.
- ISO 9000 is NOT something management can simply delegate to employees and/or a consultant.

## 4.2 Quality System

- This element requires that your quality related systems be defined within company documentation.
- You will have to define how you 'Plan' for quality

### 4.3 Contract Review

- This element requires contracts to be reviewed for a number of concerns and issues.
- The intent is to ensure everyone agrees on terms and requirements

## 4.4 Design Control

 Does not typically apply to transportation companies, but can address design if you wish to.

### 4.5 Document and Data Control

- This requires that 'quality related' documentation be controlled.
- Data (quality related forms) must also be defined and controlled.

## 4.6 Purchasing

 This element requires a company to review suppliers of material and services. Each transportation company is different, but most will have some part of this requirement to satisfy.

## 4.7 Customer Supplied Product

- This requirement originally was aimed at materials utilized in the manufacture of product, but has extended to include transportation containers and related items. The main thrust is that customer supplier items be tracked and taken care of.
- May include Customer trailers.

# 4.8 Product Identification and Traceability

• In the transportation industry, the extent of identification and traceability is company dependent/specific. The expectation is that you have a system to identify and track items in transit.

### 4.9 Process Control

 This element requires that your 'processes' be controlled. In the transportation industry your process is the conveyance of goods. Things to consider may include on-time pick-up and delivery as well as no damage to conveyed goods.

## 4.10 Inspection and Testing

 This element typically relates to manufacturing oriented industries.
 Typically transportation industries do not have 'product' to inspect and/or test.

# 4.11 Control of Inspection, Measuring and Test Equipment

- This element typically applies to equipment used to measure and/or test product. Since the transportation industry does not typically manufacture product it \*may\* not apply.
- However, if your company does any critical maintenance, it may apply to your company.

### 4.12 Inspection and Test Status

- Typically this element applies to manufactured goods. Application to the transportation industry will be strained, at best.
- However, if your drivers must check shipments for damage (an example), you could be subject to inspection

## 4.13 Control of Nonconforming Product

 While this is aimed at manufacturers, and there is technically no manufactured product, you can expect your service to be addressed. This is to say, if you deliver late, your service is nonconforming (assuming you give some assurance to the customer that you will deliver on a certain day at a certain time).

## 4.14 Corrective and Preventive Action

- This element requires you to address problems in your service by a documented methodology and that you show proof that your corrective measures are effective at eliminating the problem
- The goal is Preventive Action

# 4.15 Handling, Storage, Packaging, Preservation and Delivery

 This originally was aimed at manufactured product. In the transportation industry, depending on how your company services are structured, you may or may not have one or more of these specifics apply to you.

## 4.16 Control of Quality Records

 This element is most important. It requires that 'quality related data' be defined and controlled. Each company has to decide and document what theirs are. Disposition also has to be addressed.

## Typical Types of Records

- Contract Review
- Purchasing
- Identification and Traceability
- Process Control
- Inspection and Test
- Control of Measurement and Test Equipment
- Non-conforming Product
- Corrective and Preventive Action
- Internal Quality Audits
- Training



## Records Management Activities

- Management of Active records
- Records creation (forms)
- Design of records system
   Retention schedule
   Vital records protection
- Development of records procedures

Indexing

Filing

Access

Disposition



## 4.17 Internal Quality Audits

- This element requires that at least once a year your company audit its self against its internal documentation. This can be done by company employees or a company can sub-contract the service. However, this is in addition to the registrar's annual 'visit'
- Internal audits can be 'farmed out' (they do not have to be performed by company personnel)

### 4.18 Training

 Generally, in the transportation industry the CDL serves as evidence of proficiency. However, the company has to be ready to show that any need for training is identified and carried out.

## 4.19 Servicing

 This element is aimed at manufacturers who provide services to their customers in setting up and/or maintaining product. Transportation being a service industry this element does not strictly apply. However, many companies have people who actively support their customers.

### 4.20 Statistical Techniques

 The element requires a company to identify places where one or more statistical techniques would benefit their operations. In the transportation industry, on-time delivery would be one data point which would benefit a company in monitoring.

### ISO 9000 Reminders

- Does NOT define quality
- Is NOT a one-time process
- Is NOT easy
- Requires commitment
- Requires resources



### **Trends**

- Trend towards international agreements in business and trade (e.g.: NAFTA & GATT)
- ISO 14000: Documented environmental quality system
- Occupational Safety and Health
- Federal Transportation Agencies

### Wrap-Up

#### How To Begin

- Examine your commitment
- Examine your reasons
- Define your resources
- Further your education

### Not a panacea, cannot neglect...

- Marketing 101
- Competitive Strategy 101
- Human Resources 101
- Project Management 101

## **Customer Satisfaction**

- Improve Product
- Improve Performance
- Increase Business
- Increase Profits

### Additional Resources

Tune your browser to http://www.qs9000.com

 National ISO Support Group (616) 891-9114

 Society of Automotive Engineers (412) 776-4841

American Society for Quality

(Formerly the ASQC)

(800) 248-1946

Society of Manufacturing Engineers

(800) 733-4763