

# Quality Assurance

Contents	page
<hr/> <b>Introduction</b>	1
<hr/> <b>History and evolution of the quality discipline</b>	1
<hr/> <b>Development of quality standards</b>	1
<b>BS EN ISO 9001: 2000</b>	2
- Structure and philosophy	2
- Standard requirements	3
<hr/> <b>Quality schemes</b>	3
- Multi product/sector schemes	4
- Management systems/product conformity	4
- British Board of Agrément	4
- Kitemark	4
<hr/> <b>Elements of a quality system</b>	4
<hr/> <b>European standardisation and quality requirements</b>	5
<hr/> <b>Glossary of terms</b>	7
<hr/> <b>Self-assessment questions</b>	8
<hr/> <b>Answers to self-assessment questions</b>	9



### **Health and safety**

*All mortar mixtures, both wet and dry, are abrasive and alkaline. When working with wet mortar, waterproof or other suitable protective clothing should be worn. Guidance on the use of these materials can be found in MPA Mortar data sheet No. 20.*

## Introduction

In looking at quality assurance, this learning text covers the history of quality and the development of the quality discipline together with the creation of the function of dedicated quality personnel. It also covers the development of quality standards, the various types of assessment and quality schemes and systems that exist.

The text also looks at quality assurance and its relevance in relation to European standardization, including CE marking. A glossary of terminology and self-assessment questions are also included.

## History and evolution of the quality discipline

Quality assurance is not a recent innovation. Quality rules existed at the time of the Roman Empire. Emperor Quintus instructed that goods suppliers to the imperial and senatorial households had to affix their mark to their goods. If these failed in service the manufacturer was likely to be punished.

In the UK, early mention of quality assurance dates from the reign of Edward I when in 1300, legislation on jewellery hallmarking was introduced. A further development in 1340 was legislation creating the role of clerks in markets, whose job was to check weights and measures. Medieval trade guilds acted as controllers of quality, laying down regulations for the training of apprentices, the behaviour of members of the guild and price regulation. A famous individual who worked in the quality field was Geoffrey Chaucer who for a period was employed as a quality representative to the royal armoury.

In early times when the skilled craftsman was at the pinnacle of the workforce, he would make an object, a piece of furniture or some other item. At each stage production, he would examine it to ensure that it was an acceptable piece of work according to standards determined by himself. A second, essentially identical item produced after the first, would probably not be exactly the same. It would nevertheless have the same known skills of the craftsman embedded in it and would have been produced to the same

high standards which the craftsman imposed on himself.

The decline of the agricultural society and the emergence of manufacturing industry, initially as small production units, created new problems. Initially factories produced complete products or sub assemblies, which were either accepted or rejected by customers.

Gradually, two factors necessitated the creation of a culture of more rigorous inspection:

- Advances in engineering demanded production to greater dimensional accuracy
- The increasing tendency to manufacture components separately and then assemble them elsewhere.

These developments created the need for inspection personnel who initially reported to the production department. However, conflicts of interest soon arose and in some cases separate inspection departments were set up.

Impetus for many changes in society has been war. David Lloyd George, before becoming prime minister was minister of war. He was concerned that more planes fell from the sky during World War One because of electrical and mechanical failure than were shot down. His concern led to the introduction of inspection systems into military procurement.

The establishment of the process of inspection was an acknowledgement that the production process was not as efficient as it might have been. If its primary purpose was to ensure that sub-standard goods never reach the customer then it soon became apparent that the objective had not been achieved. The creation of inspection departments brought a number of new issues into prominence such as the recording of data, the precision of measuring instruments and the introduction of uniform standards. It soon became apparent that in addition to product inspection, other duties were required and gradually quality control departments evolved.

In the early part of the 20th century, Fredrick

W Taylor, chief engineer of the Mid Vale steel works in the USA introduced the concept of scientific management. Around the same time, statistical theory began to be applied to quality control. One of the first individuals to apply the new methods was Walter A Shewhart of the Bell Telephone company. In 1924 Shewhart put forward the idea of a control chart and in 1931 he published a book entitled "Economic control of quality of manufactured products".

In the UK, the Institution of Engineering Inspection was set up in November 1922. This organisation was open to all those engaged in inspection activities both within the public and private sectors. The Second World War led to a dramatic rise in demand for munitions and there simply was not sufficient time to check that they were manufactured to the desired quality. It therefore became very important that quality was built into the manufacturing process. Inspection had been the system used by almost all manufacturers to prevent unacceptable products from reaching their customers. Gradually it was realised that the implementation of a system that detected faults as they occurred was a more efficient method to prevent those faults occurring in the first place.

Following the Second World War, the industrial system in Japan had been virtually destroyed. Additionally, Japan had acquired a reputation for producing cheap imitation products. The Japanese recognised their problems and started to take action to resolve them. They received assistance in quality improvement from the civil communication section of the occupation forces. Further assistance was provided by one of the leading figures in American quality management Dr W Edwards Deming, who had worked very closely with Shewhart. Over a number of years adoption of the systems put forward by Deming transformed Japanese industry.

## Development of quality standards

Formalised quality requirements were introduced by the American military forces with a series of written standards. Mil-Q-9858 was published in 1959 and covered quality control systems for industry. Other standards

dealt with the subject of inspection (eg, Mil-I-45208A). The creation of the North Atlantic Treaty Organization (NATO) led to the concept of formal quality standards moving from a national to an international arena. NATO published a series of quality standards entitled Allied Quality Assurance Publications (AQAPs) in 1968.

Colonel Rabey of the British defence establishment advocated a fundamental change in the way that the Ministry of Defence (MOD) approached its suppliers. He recommended that the supplier be made responsible for the quality of product or service delivered, that the technical competence of potential suppliers was assessed before contracts were awarded and that the ministry should ensure that suppliers had effective systems in place. The recommendations put forward led to the introduction of a series of quality standards (the 05 series) being introduced across all MOD contracts.

The success of the 05 series led to the publication of British Standard BS 5750: Quality Systems, which was first published in 1979. This was revised in 1987. Increasing international interest in quality standards led to ISO (the International Standards Organisation) publishing the ISO 9000 series of standards in 1987 and updating and re-issuing them in 1994. CEN (Comite Europeen De Normalisation) had also published a series of quality standards as the EN 29000 series but these were withdrawn and the requirements incorporated into the 9000 series. The British Standards Institution (BSI) also withdrew BS 5750 and adopted the International Standard as BS EN ISO 9001. This standard was in three parts:

BS EN ISO 9001:1994 Quality Systems – Model for quality assurance in design, development, production, installation and servicing.

BS EN ISO 9002:1994 Quality Systems – Model for quality assurance in production, installation and servicing.

BS EN ISO 9003:1994 Quality Systems – Model for quality assurance in final inspection and test.

Organisations selected the part that was most for them. In addition to the three parts of the

standard, which prescribed the requirements, other parts, which provided guidelines on their implementation were also published.

## BS EN ISO 9001:2000

### Structure and philosophy

The year 2000 version of the quality standard has the requirements in a single standard and there is no longer a 9002 and 9003. The standard is now composed of the following:

BS EN ISO 9000:2000 – Quality management systems – Fundamentals and vocabulary  
BS EN ISO 9001:2000 – Quality management systems – Requirements

BS EN ISO 9004:2000 – Quality management systems – Guidelines for performance improvement

BS EN ISO 9000:2000 contains the vocabulary and the underlying philosophy of the standard. This document is descriptive and although it does not contain any mandatory material, it does contain a description of eight key management principles.

BS EN ISO 9001:2000 is the standard against which an organization will be audited and certificated.

BS EN ISO 9004:2000 has been prepared to give guidance on continuous improvement within the organization.

The standard (BS EN ISO 9001) has been structured in to eight main clauses, these being:

- Scope
- Normative reference
- Terms and definitions
- Quality management system
- Management responsibility
- Resource management
- Product realization
- Measurement, analysis and improvement.

The main requirements are contained within clauses 4 to 8. The standard has adopted a process approach rather than the procedure based approach of the 1994 version. The year 2000 version has been based on eight quality management principles, these being:

- a) A customer focused organisation. The

writers of the standard took the view that companies or organisations depend on their customers and therefore they should understand their current and future needs. This objective also recognises that companies should meet customer requirements and make efforts to exceed customer expectations.

b) Leadership. Emphasis has been placed on leaders establishing unity of purpose, direction and the internal working environment of the organisation (company). The standard has been drafted to take account of the view that leaders can create the environment in which individuals can become fully involved in achieving the organisation's objectives.

c) Involvement of people. The standard recognises that people at all levels are the essence or lifeblood of a company and that it is important to harness their abilities for the company's benefit.

d) A process approach. This recognises that the desired result can be achieved more efficiently when related resources and activities are managed as a process.

e) A system approach to management. This involves identifying, understanding and managing a system of interrelated processes for a stated objective contributing to the efficiency and effectiveness of a company.

f) Continual improvement. The standard recognises that continual improvement is a prime requirement.

g) A factual approach to decision-making. This acknowledges that effective decisions are based on logical analysis of data and information.

h) A mutually beneficial relationship. This is based on the philosophy that the ability of a company and its suppliers to create value is embraced by mutually beneficial relationships.

Perhaps the most radical change in the year 2000 version of BS EN ISO 9001 is the process approach. A process is a sequence of related tasks with inputs and outputs. A process can also be represented as a loop, which emphasises the interlinking of

the component parts of a process. In any organisation, there are three principal types of approach:

- A procedure-based organisation focuses on doing the right things, although not necessarily based on stakeholders' needs.
- A process-based organisation focuses on doing the right things based on stakeholder needs.
- A process-based organisation with appropriate procedures, focuses on doing the right things correctly, in line with stakeholders' needs.

#### Standard requirements

The quality standard BS EN ISO 9001:2000 requires that a documented system be established. Clause 4.1 states "The organisation shall establish, document, implement and maintain a quality management system." The standard requires that, as a minimum, documentation is required in respect of just six procedures:

- Document control
- Record keeping
- Non-conformance control
- Auditing
- Corrective action
- Preventative action.

Company management must decide in all other cases whether it needs additional information to facilitate control. It is normal practice to maintain the quality system requirements within the quality manual. Clause 4.2.4 requires that records be maintained to provide evidence of the effective operation of the quality management system. Clause 5.3 which is entitled "Quality policy" requires that:

- The quality policy is appropriate to the purposes of the organisation.
- A commitment to comply with requirements and continually improve the effectiveness of the quality management system is stated.
- A framework for establishing and reviewing quality objectives is implemented.

- The quality policy is communicated within the organisation.
- The quality policy is reviewed and updated.

Clause 5.6 of the standard prescribes the management review, a prime requirement of any quality system. This is generally undertaken on an annual basis and should examine all aspects of the quality scheme, (eg, audit reports, customer complaints, preventative and corrective action requests and areas for potential improvements). Clause 5.6.2 of the standard lists the items that should be considered during the management review.

Clause 8.5.2 of BS EN ISO 9001 states "the organisation shall take corrective action to eliminate the cause of the nonconformities in order to prevent recurrence".

Clause 8.5.3 of BS EN ISO 9001 states "the organisation shall identify preventative action to eliminate the causes of potential non-conformities in order to prevent their occurrence". Thorough auditing frequently identifies potential non-conformities.

The two terms corrective and preventative action sometimes cause confusion. A corrective action is essentially a backward looking phenomenon, which identifies the last point from which action can be taken to rectify a mistake. Preventative action is essentially a forward-looking process to determine at the earliest point what action is necessary in order to prevent an error occurring.

#### BS EN ISO 9001:2008

ISO 9001:2008 basically restates ISO 9001:2000. The 2008 version only introduced clarifications to the existing requirements of ISO 9001:2000 and some changes intended to improve consistency with ISO 14001:2004. There were no new requirements. For example, in ISO 9001:2008, a quality management system being upgraded just needs to be checked to see if it is following the clarifications introduced in the amended version.

#### Forthcoming 2015 version

The next version of the standard is expected to be published in December 2015.

## Quality Schemes

Quality assessment may be classified as first, second or third party: First party assessment involves an organisation setting up a quality system, documenting the structure and procedures and subsequently notifying customers that a quality system has been implemented.

Second party assessment involves documenting a quality system and inviting the customer to collaborate in its development for quality as specified by the customer. Alternatively a client or customer may undertake its own assessment of an organisation's quality assurance scheme. Historically second party assessment was undertaken in the construction industry by consulting engineers or clerks of works acting on their behalf.

Third party assessment involves an organisation independent of the producer or purchaser (customer) undertaking the assessment process.

The number of third party quality assurance schemes grew in the late 1970s and early 1980s. Central government was concerned that the lack of uniformity of standards between the various schemes was leading the certification industry into disrepute. This was also having an adverse effect on the reputation of UK industry in the international trading market. This led to a government white paper, "Standards, Quality and International Competitiveness", being published in July 1982.

In 1984 the Department of Trade and Industry set up the National Accreditation Council for Certification Bodies (NACCB) under a memorandum of understanding with BSI. The NACCB was empowered to grant recognition to certification bodies who reached a prescribed minimum standard. These bodies were given the privilege of using the symbol of the crown and the gold tick of approval on their stationary. The mark symbolising approval was also permitted to be used by companies whose quality scheme was approved by a NACCB certification body. Once a certification body was accredited by NACCB, it was subject to regular surveillance visits to ensure standards were maintained.

In addition to a regulator for certification bodies, the government also established under the auspices of the National Physical Laboratory, schemes for the certification of testing (NATLAS) and calibration laboratories (NAMAS). In 1985 these two organisations were merged under the NAMAS banner. Central government commissioned a report from independent management consultants on the future of UK certification activities. This led in 1995 to the creation of a single overseeing body for all testing and certification activities – the United Kingdom Accreditation Service (UKAS). However, a number of independent certification bodies which are not recognised by UKAS still continue to operate.

UKAS now operates as a company limited by guarantee under a memorandum of understanding with the Department of Trade and Industry. A number of industry panels have been created to advise UKAS staff on the needs of different sectors of industry.

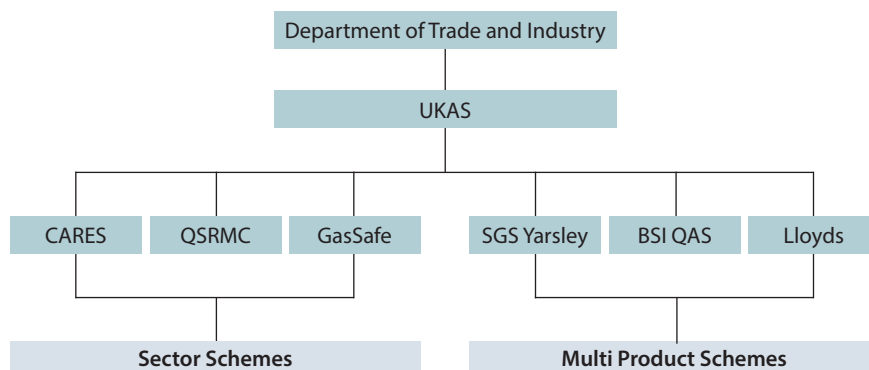
#### Multi product/sector schemes

Certification bodies can be categorised under two headings:

- Certification under a registered industry sector scheme. Some industries have set up quality assurance organisations which have been accredited by UKAS to provide certification for a particular industry sector.
- Direct certification. This is where an organisation approaches an independent certification body directly for third party certification. There are a number of sector schemes; two that are especially relevant to the construction industry are CARES (the Certification Authority for Reinforcing Steels) and QSRMC (the Quality Scheme for Ready Mixed Concrete). Other well known schemes include GasSafe which is applicable to the installation of gas appliances.

Sector schemes only offer certification in a limited range or even only for a single product. Multi-product organisations (eg, British Standards Institution Quality Assurance Services, Lloyds Register Quality Assurance, SGS Yarsley) offer certification across a diverse

Figure 1: **The organization of Quality Schemes**



range of products, often on a global basis. A number of the multi product schemes also operate sector schemes, eg, the cement industry certification scheme operated by BSI-QAS.

#### Management systems / product conformity

Some organisations do not have a physical product to test (eg, law firms, educational establishments) but nevertheless wish to have their management systems certified by a third party quality assurance organisation. Other organisations only elect to have their management systems accredited by a third party. This is known as Category 1 certification and is often a transitional step prior to seeking product conformity certification, known as Category 2, not only provides certification that the quality management system is operating in accordance with the required standard, but that the product is also complying with the appropriate quality standard.

#### British Board of Agrément

The British Board of Agrément (BBA) was established in 1966. It was originally called the Agrément Board, the present name being adopted in 1982. It is involved in the testing, assessment and certification of products used within the building and construction industry. The types of product assessed are wide ranging but are generally characterised by being new or innovative, although existing products may also be assessed.

The BBA is an independent company limited by guarantee and is controlled by a governing board nominated by the government. It is

based at the Building Research Establishment near Watford in Hertfordshire.

An Agrément certificate provides assurance that a product or system to which the certificate relates, if properly used in accordance with the terms of the certificate, will meet the relevant requirements. Some publications refer to BBA approval as Category 3 certification.

#### The Kitemark

British Standards Institution was set up in 1901 and has operated a certification scheme since 1903: the first British Standard Mark was granted for tramway rails in 1903. The Kitemark symbol was adopted in 1922 and the first licence granted to the General Electric Company in 1926. A licence to use the Kitemark can be obtained by a manufacturer who can demonstrate to the satisfaction of BSI that he can consistently manufacture the product in accordance with the requirements specified in the appropriate standard. The capability of the manufacturer is assessed by evaluating its quality system against the requirements of BS EN ISO 9001 and testing the product against the requirements of the product standard. Surveillance of the quality system and the product is also undertaken on an ongoing basis.

## Elements of a quality system

The section earlier on the history and evolution of the quality discipline stated

the importance of building quality into the manufacturing process. To ensure a quality product or service all those involved must:

- Be aware of what is to be done which may involve having the appropriate standards, specifications and, where appropriate, drawings.
- Be aware of how the task is to be performed, which involves the provision of suitable training and the availability of appropriate work instructions.
- Possess the necessary resources, plant and materials.
- Be aware if the task has been properly completed, which may involve inspection, measurement or testing of a product.
- Possess suitable motivation to accomplish the task at the required quality level.
- Maintain appropriate records.

## European standardization and quality requirements

The change from national to European standards has increased the necessity to implement quality assurance. For some products third party quality assurance is mandatory. Now that the European Union is determined to prevent barriers to trade, national standards are believed to be a major obstacle to the free movement of goods and services. In different industrial sectors legislative statements, known as directives, have been introduced to attempt to remedy this situation. The change to European Standards has resulted in some new terminology being introduced (eg, mandate, harmonised standard, etc). Definitions of these terms are given in the glossary of terms included in this learning text. The Construction Products Directive (CPD) has the objective of breaking down barriers to trade within the design, materials and component supply industries of the construction sector. To achieve this the CPD has a number of elements:

- A system of harmonised technical specifications.
- The CE marking of products (to demonstrate product conformity).
- A system of Notified Bodies (to administer CE marking).

To comply with the requirements of the CPD, a product subject to a mandate and placed on the market has to comply with the following essential requirements:

- Mechanical strength and stability.
- Security in the case of fire.
- Health, safety and the environment.
- Security for user.
- Noise protection.
- Energy economy and thermal insulation.

A product that meets the essential requirements of a harmonised standard may be placed on the market. It is a requirement that conformity to the standard is indicated by the affixing of the CE mark to the product, to its packaging or to the commercial documentation where there is no packaging. The CE mark is not a quality mark: it only

Figure 2



signifies that the essential requirements of the CPD have been met. The CE mark may be regarded as a passport to assist in the movement of goods within the European Union. Figure 2 shows an example of the CE mark.

It is important to note that where a mandate has been issued, only products that comply with the requirements of the technical specifications (ie, standards) may be placed on the market.

Not all products are regulated. In order for a harmonised standard to be produced the European Commission must issue a mandate to CEN. A mandate is an instruction given by the European Commission to CEN to prepare a harmonised standard.

Where such a harmonised standard has been issued, EU member countries are required to withdraw conflicting national standards within a given time period. The time period may vary because some product standards are dependent on supporting testing standards being available.

Normally CEN identifies the supporting standards and groups these into packages. The date of withdrawal of conflicting national standards is generally fixed at a number of months after the last standard in the package becomes available.

Within a harmonized standard not all requirements are covered by the mandate - normally the essential requirements are listed in an "Annex ZA" to the standard. The process of judging whether a product meets the requirements for the affixing of a CE mark is called attestation of conformity.

Not all products are judged at the same level. Each harmonised product standard prescribes the level for judging attestation of conformity. The levels range from self-certification (level 4) to third party product testing (level 1+) as well as the operation of a third party quality assurance system. The levels of attestation of conformity that have been set for mortar, screed and their constituent products are:

**Cement:** level 1+

**Admixtures:** level 2+

**Aggregates:** level 4

**Mortars:** level 2+ (designed masonry mortars)  
level 4 (other masonry mortars and rendering mortars)

**Screeds:** level 4 (for some applications)  
level 3 may be required)

Table 1 (overleaf) lists the requirements for each of the levels:

Table 1: **Attestation of Conformity Systems** (see the Glossary for further information on terminology)

Conformity numbering system	1+	1	2+	2	3	4
<b>Tasks for the manufacturer</b>						
Factory production control	✓	✓	✓	✓	✓	✓
Extra testing of samples taken from the production unit in accordance with a prescribed test plan	✓	✓	✓	•	•	•
Initial type testing	•	•	✓	✓	•	✓
<b>Tasks for the notified body</b>						
Initial type testing	✓	✓	•	•	✓	•
Initial certification of factory production control	✓	✓	✓	✓	•	•
Continuous surveillance of factory production control	✓	✓	✓	•	•	•
Audit testing of samples taken from the factory or the market place or site	✓	•	•	•	•	•

The monitoring of CE marking in each country is regulated by the Notified Bodies (which may be some of the quality assurance organizations).

In addition to harmonised standards, voluntary standards also exist. This means that the standard does not have the force of European law and a product covered by a voluntary standard cannot be CE marked. However under an agreement between the national standards bodies, conflicting national standards will have to be withdrawn.

# Glossary of Terms

## **Attestation of conformity**

A declaration required by Chapter V of the CPD, attesting that a product meets the requirements of a harmonised European Standard. The process of testing and quality assurance to achieve this is called the evaluation of conformity.

## **BSI**

British Standards Institution

## **CEN**

European organisation for standardisation and the abbreviation for Comité Européen De Normalisation.

## **CE marking**

Signification that a product complies with the essential requirements of the CPD and with a harmonised European technical specification or standard.

## **Class**

A combination of two levels between which the performance must fall: (eg, strength between 32.5 – 52.5 N/mm<sub>2</sub>)

## **Compliance**

Compliance with the standards - an area of confusion exists as to what has to be complied with. There are two levels of compliance:

- Compliance with the voluntary parts will be necessary to claim one is supplying in accordance with the standard. This will be necessary for commercial reasons.
- Compliance with the harmonised part will be necessary to comply with the law.

## **CPD**

Construction Products Directive (published by the European Commission in 1988; incorporated into European Law in 1989 and passed into UK law by the Construction Products Regulations, 1991). Potentially, it applies to all products produced for permanent incorporation in buildings and civil engineering works. A directive is a legal device used by the European Union to establish policy at European level.

## **Harmonised Standard**

A harmonised standard has the objective of satisfying the essential requirements of a directive. Harmonised standards define the relationships between national regulators and producers. The national regulators (normally UKAS in the UK), will delegate powers to notified bodies (third party quality assurance bodies, eg, British Standards Institution: Quality Assurance Services), to carry out much of the work. Some standards will not have a mandate and will therefore only satisfy a need between the producer and the purchaser. A standard that does not have a mandate cannot be harmonised and the product cannot carry a CE mark. This type of standard is called a "voluntary standard". Once a harmonised standard is published, conflicting national standards have to be withdrawn within a set time period. Any product that satisfies the CPD and has a CE mark must be allowed to be placed on the market in all European Union countries.

## **Level**

Performance which will be exceeded: eg, strength greater than 32.5N/mm<sub>2</sub>.

## **Mandate**

A political request from the European Union to CEN to produce a standard in a specific product area. The issuing of a mandate leads to the production of a harmonised standard.

## **Voluntary standard**

A voluntary standard is produced by CEN, normally to assist in free trade. Under CEN rules, national standard bodies, eg, BSI, have to withdraw conflicting national standards. Concrete for instance, has not yet been granted a mandate and therefore, the concrete Standard EN 206-1 cannot be harmonised and no CE mark is applicable.

*Note: Most standards are prepared at the request of industry, but the European Commission may also request that some standards are prepared to implement European legislation.*



---

## Self-assessment questions

1 What is the number and title of the main quality standard in use in the UK?

A \_\_\_\_\_

2 When was BS 5750 first published?

A \_\_\_\_\_

3 Designed masonry mortar requires level 2+ attestation of conformity, what are the requirements involved?

A \_\_\_\_\_

\_\_\_\_\_

4 What item of European legislation governs the CE marking of mortar?

A \_\_\_\_\_

5 What are the eight main clause headings in BS EN ISO 9001:2000?

A \_\_\_\_\_

\_\_\_\_\_

6 What is the role of UKAS?

A \_\_\_\_\_

7 The year 2000 version of BS EN ISO 9001 requires as a minimum procedures in respect of six activities - what are they?

A \_\_\_\_\_

\_\_\_\_\_

8 What is the difference between a category 1 and a category 2 quality scheme?

A \_\_\_\_\_

\_\_\_\_\_

9 When was the first BSI kitemark awarded?

A \_\_\_\_\_

10 What is the difference between preventative and corrective actions?

A \_\_\_\_\_

\_\_\_\_\_

## Answers to self-assessment questions

- 1 BS EN ISO 9001:2000 Quality management systems – Requirements.
- 2 1979.
- 3 The implementation of a factory control system, initial type testing and the taking of samples in accordance with a prescribed plan. Additionally the notified body has to certify the factory production control system and undertake continuous surveillance of the system.
- 4 The Construction Products Directive.
- 5 Scope, Normative references, Terms and definitions, Quality management system, Management responsibility, Resource management, Product realisation and Measurement analysis and improvement.
- 6 The accreditation of certification bodies and laboratories.
- 7 Document control, record keeping, nonconformance control, auditing, corrective action and preventative action.
- 8 A category 1 scheme only covers management systems, a category 2 scheme covers management systems and product conformity.
- 9 1926.
- 10 Preventative action is intended to eliminate the occurrence of potential non-conformities. Corrective action is taken to eliminate the recurrence of nonconformities.

## MPA Mortar Learning Texts include:

- 1 Introduction to modern mortars
- 2 Cementitious materials
- 3 Aggregates
- 4 Admixtures, additives and water
- 5 Brick and block production
- 6 Properties of masonry mortar
- 7 Production, delivery and storage of mortar
- 8 Mortar testing
- 9 Specifications
- 10 Quality assurance
- 11 Construction
- 12 Properties of rendering mortar
- 13 Best practice - potential site problems



MPA Mortar is part of the Mineral Products Association, the trade association for the aggregates, asphalt, cement, concrete, dimension stone, lime, mortar and industrial sand industries.

**Mineral Products Association Ltd**  
First Floor  
297 Euston Road  
London NW1 3AD  
Tel 0203 978 3400  
mick.russel@mineralproducts.org  
www.mortar.org.uk

Factory produced mortar products will contain either cement or lime, both of which have properties which are hazardous to health. Please refer to the manufacturers or suppliers Material Safety Data Sheet for the specific product/grade to find more information on the nature of the hazardous properties, the risks and health effects of exposure and the recommended safe use and handling procedures.