

Changes to the PPE Directive

The history and future of PPE Certi cation

A whitepaper



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at a glance

The Personal Protective Equipment (PPE) Directive 89/686/EEC is a fundamental piece of European legislation relating to occupational safety throughout Europe. It can be described as covering any device or appliance designed to be worn or held by an individual for protection against one or more health and safety hazards, and it is mandatory if a PPE product falls within the scope of the directive. However, there are some exemptions such as PPE used by the military and for law enforcement.

The PPE Directive was rst adopted by the European Council on 21st December 1989. It was implemented into UK law as the Personal Protective Equipment (EC Directive) Regulations 1992 (SI 1992/3139) and known as the 'Principal Regulations'. These Regulations were made on 10th December 1992 and came into effect on 1st January 1993.

Prior to European EN (European Norm) speci cations and CE marking, individual states produced standards for PPE. As the need for standardisation grew across Europe, EN standards started to be written. These standards are known as Harmonised Standards and are listed in a document called the Of cial Journal (OJ) which is available online at:

The PPE Directive was one of the rst New Approach Directives and is now over 20 years old. In order to re ect current technologies and processes for developing and bringing PPE to the market, it needs to be updated.

It is also important to bring it in line with other Directives that have undergone

also mean that the old Directive will rather than remain in its current status. This means that the new Regulation will not have to be transposed into each Member State's national law. (A Directive is a legislative act that sets out legislation.)

a revision in recent years. The changes an objective that all EU countries must achieve by a given date. However, it is up now be re-implemented as a Regulation to the individual countries to decide how this is done. In contrast a Regulation is a binding legislative act, and it must be applied in its entirety across the EU without the need for separate national

What are the

The draft version of the Regulation has already been approved by the European Commission and Parliament. The next phase involves the European Council to reach an agreement on the text by the end of summer 2015. There are a number of changes that are being proposed taking place including:

Moving hearing protection from Category 2 to Category 3 PPE

Changing life jackets from Category 2 to Category 3 PPE Issuing a Declaration of Conformity with each PPE or at least a link to where it can be obtained

?

Possibly covering domestic PPE (for example oven gloves)

Bringing the Regulation in line with similar European requirements such as the Medical Devices Directive by suggesting a ve-year certicate validity

Once the current Directive is re-issued as a Regulation in 2018, the scope is likely to be changed signi cantly. The current proposed wording for this is described in the proposed scope below.

The

It is currently proposed that the following products fall under the scope of the Regulation. This is important as it will help you to understand whether your products are covered by the Regulation.

- This Regulation shall apply to personal protective equipment (PPE)
- 2 This Regulation shall not apply to PPE that is:
 - speci cally designed for use by the armed forces or in the maintenance of law and order:

of the Regulation

designed to be used for selfdefence with the exception of PPE intended for sporting activities;

designed and intended for private use to protect against atmospheric conditions that are not of an extreme nature:

for exclusive use on seagoing vessels or aircraft that are subject to the relevant international treaties applicable in Member States; for head, face or eye protection of users, subject to Regulation 22 of the United Nations Economic Commission for Europe (UNECE) on uniform provisions concerning the approval of protective helmets and of their visors for drivers and passengers of motor cycles and mopeds.

Where to from here?

All manufacturers of PPE need to be aware that any existing certifications they currently hold will expire when the Regulation comes into force at the end of 2018. So it is therefore important to keep up to date with the coming changes and prepare for the impact on business. This would also apply to new manufacturing and distributor companies.

At this stage you need to consider the following:-

Because new EC Type Examination certi cates will have to reference the speci cs, all products currently certi ed to

old withdrawn standards will have to be tested to the latest versions.

- Look at existing product ranges and ensure that they are to the latest product speci cations
- If you are placing products that will change category such as life jackets or hearing protection etc. onto the market, be aware of what the change in classi cation will mean.
- If you are a distributer, be aware that there are parts of the directive that will have direct implications to you. As more information becomes available we will produce updates on this.

Obligations for

manufacturers

The PPE Regulations are mandatory covering any type of product that falls within its scope. If you are therefore in the PPE industry, it is a legal requirementmeans when the regulation comes into to comply. Previously the PPE Directive force that distributors or anyone in the focused on manufacturers placing

products onto the market, but when the new Regulation becomes effective the whole supply chain will be involved. This and that they make available on the supply and distribution chain should

take appropriate measures to ensure that PPE meets standard requirements market only products which comply with the Regulation.

Explaining the

Category 1 – Simple PPE

PPE in this category is designed to protect users against minimal risks. These include as examples:

super cial mechanical injury;

contact with water or cleaning materials of weak action;

contact with hot surfaces not exceeding 50°C;

damage to the eyes due to exposure to sunlight (other than during observation of the sun);

atmospheric conditions that are not of an extreme nature.

Category 2 – Intermediate PPE

PPE in this category protects users against risks other than those listed in categories I and III, as well as made-to-measure PPE (excluding PPE that falls under category I. The following products are included as examples:

Safety spectacles and goggles Industrial helmets and bump caps Bicycle helmets

of PPE

Category 3 – Complex PPE

PPE falling under this category is designed to protect against mortal danger or dangers that may seriously and irreversibly harm health; the immediate effects of which the designer assumes that the end user cannot identify in suf cient time.' Risks include:

inhalation of harmful substances;

aggressive chemicals;

ionising radiation;

high-temperature environments the effects of which are comparable to those of an air temperature of at least 100°C; low-temperature environments the effects of which are

comparable to those of an air temperature of -50°C or less;

falling from a height;

electric shock and live working;

drowning;

cuts by hand-held chain-saws;

- high-pressure cutting;
- bullet wounds or knife stabs; harmful noise.

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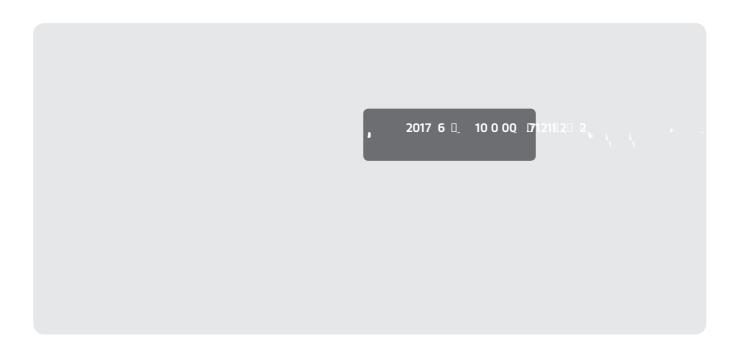
L ki g at your existing product ranges and identifying those that are tested to withdrawn standards. Start to prepare to get them tested to the latest speci cations.



Thi ki g about how you will meet the change in classi cation if you place hearing protection or life jackets onto the market. The BSI Kitemark would ensure that you meet all your ongoing surveillance requirements, both to the current PPE Directive and the new regulations when they come into force.

ac i g us to check how you will comply to the new Regulation if you are a manufacturer or distributor of products like dish-washing gloves and oven gloves.

Timelines



Support from.

How BSI can help you through the changes.

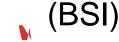
We will continue to monitor the proposed changes and keep all our clients updated as more information is released from the European Commission. This information will be available on our website and we will be sending out regular updates, whitepapers and offering training courses. Our Certi cation

Managers, Test Engineers, and other colleagues will be kept up to date on the changes so that we can let you know in advance about the impact that these changes may have.

BSI is a Noti ed Body - look at our scope of PPE Standards http://www.ukas.org/Certi cationBodies/schedules/ PROD/0003Product%20Certi cation.pdf

The role of a

BSI is a Noti ed Body for the PPE Directive and numerous other EU Directives. We have a comprehensive scope, being able to carry out Article 10 (EC Type Examination), Article 11A (on-going surveillance through annual testing) and Article 11B (on-going surveillance through annual factory) for head protection, eye protection, clothing,



gloves, respirators, footwear and life jackets as well as many other devices. We can issue EC Type examination certi cates as well as carry out on-going surveillance globally, delivered locally where at all possible.

In the UK a Noti ed Body is a body which has been appointed by the department for Business Innovations and Skills

(BIS), to carry out one or more of the conformity assessment procedures to Article 10, 11A or 11B of the PPE Directive. The details of all Noti ed Bodies and their scope of approval is listed on the Europe website.

Why.

Our knowledge, expertise and drive can make a difference to your business:

We are the pioneers of many of the world's rst standards.

We are renowned for our innovative work in many elds.

We continue to lead the way with ongoing developments in various industries.

You can enjoy the bene ts of working with BSI teams who have decades of experience helping businesses of all sizes.

We provide end-to-end support, helping you monitor and maintain your excellence throughout.

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