

**DIRECTIVE 2006/121/EC OF THE EUROPEAN PARLIAMENT  
AND OF THE COUNCIL**

**of 18 December 2006**

**amending Council Directive 67/548/EEC  
on the approximation of laws, regulations and administrative  
provisions relating to the classification, packaging and labelling of  
dangerous substances in order to adapt it to Regulation (EC) No 1907/2006  
concerning the Registration, Evaluation, Authorisation and Restriction  
of Chemicals (REACH) and establishing a European Chemicals Agency**

THE EUROPEAN PARLIAMENT AND THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty establishing the European Community, and in particular Article 95 thereof,

Having regard to the proposal from the Commission,

Having regard to the Opinion of the European Economic and Social Committee<sup>1</sup>,

After consulting the Committee of the Regions,

Acting in accordance with the procedure laid down in Article 251 of the Treaty<sup>2</sup>,

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<sup>1</sup> OJ C 294, 25.11.2005, p. 38.

<sup>2</sup> Opinion of the European Parliament of 17 November 2005 (OJ C 280 E, 18.11.2006, p. 440), Council Common Position of 27 June 2006 (OJ C 276 E, 14.11.2006, p. 252) and Position of the European Parliament of 13 December 2006 (not yet published in the Official Journal).

Whereas:

In view of the adoption of Regulation (EC) No 1907/2006<sup>1</sup>, Directive 67/548/EEC<sup>2</sup> should be adapted and its rules on the notification and risk assessment of chemicals deleted,

HAVE ADOPTED THIS DIRECTIVE:

*Article 1*

Directive 67/548/EEC is hereby amended as follows:

- 1) in Article 1, paragraph 1, points (a), (b) and (c) shall be deleted;
- 2) in Article 2, paragraph 1, points (c), (d), (f) and (g) shall be deleted;
- 3) Article 3 shall be replaced by the following:

*"Article 3*

*Testing and assessment of the properties of substances*

Tests on substances carried out within the framework of this Directive shall be conducted according to the requirements of Article 13 of Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH) and establishing a European Chemicals Agency\* .

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\* OJ L"; 396, 30.12.2006, p. 1.

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<sup>1</sup> See page 1 of this Official Journal.

<sup>2</sup> OJ 196, 16.8.1967, p. 1. Directive as last amended by Commission Directive 2004/73/EC (OJ L 152, 30.4.2004, p. 1). Corrected in OJ L 216, 16.6.2004, p. 3.

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- 4) Article 5 shall be amended as follows:
- (a) in paragraph 1, the first subparagraph shall be replaced by the following:
- "Member States shall take all the necessary measures to ensure that substances cannot be placed on the market on their own or in preparations unless they have been packaged and labelled in accordance with Articles 22 to 25 of this Directive and with the criteria in Annex VI of this Directive, and, for registered substances, in accordance with the information obtained through the application of Articles 12 and 13 of Regulation (EC) No 1907/2006, save in the case of preparations where provisions exist in other Directives.";
- (b) paragraph 2 shall be replaced by the following:
- "2. The measures referred to in the first subparagraph of paragraph 1 shall apply until the substance is listed in Annex I or until a decision not to list it has been taken in accordance with the procedure laid down in Article 29.";
- 5) Articles 7 to 15 shall be deleted;
- 6) Article 16 shall be deleted;
- 7) Articles 17 to 20 shall be deleted;
- 8) Article 27 shall be deleted;

9) Article 32 shall be replaced by the following:

*"Article 32*

*References*

References to Annexes VII A, VII B, VII C, VII D and VIII of this Directive shall be construed as references to the corresponding Annexes VI, VII, VIII, IX, X and XI of Regulation (EC) No 1907/2006.";

10) Annex V shall be deleted;

11) Annex VI shall be amended as follows:

- (a) in sections 1.6.2, 1.7.2, 1.7.3, 2.1, 2.2.1, 2.2.2, 2.2.2.1, 2.2.3, 2.2.4, 2.2.5, 3.1.1, 3.1.5.1, 3.1.5.2, 3.2.1, 3.2.2, 3.2.3, 3.2.5, 3.2.6.1, 3.2.6.2, 3.2.7.2, 4.2.3.3, 5.1.3, 9.1.1.1, 9.1.1.2, 9.3 and 9.5 of this Annex, the words "Annex V" and "Annex V to this Directive", shall be replaced by the words "the Commission Regulation on test methods as specified in Article 13(2) of Regulation (EC) No 1907/2006";

(b) section 1.6.1, point (a) shall be replaced by the following:

"(a) as regards substances for which the information specified in Annexes VI, VII and VIII of Regulation (EC) No 1907/2006 is required, most of the necessary data for classification and labelling appear in the base set. This classification and labelling must be reviewed, if necessary, when further information is available (Annexes IX and X of Regulation (EC) No 1907/2006);";

(c) in section 5.1, the second paragraph shall be replaced by the following:

"The criteria set out below follow directly from the test methods set out in the Commission Regulation on test methods as specified in Article 13(2) of Regulation (EC) No 1907/2006 in so far as they are mentioned. The test methods required for the base set referred to in Annexes VII and VIII of Regulation (EC) No 1907/2006 are limited and the information derived from them may be insufficient for an appropriate classification. Classification may require additional data derived from Annexes IX or X of Regulation (EC) No 1907/2006 or other equivalent studies. Furthermore, classified substances may be subject to review in the light of other new data.";

- (d) in section 5.2.1.2, second paragraph, the second sentence shall be replaced by the following:

"Such additional scientific evidence should normally be based on the studies required by Annex IX of Regulation (EC) No 1907/2006, or studies of equivalent value, and could include:";

- 12) Annexes VII A, VII B, VII C, VII D and VIII shall be deleted.

### *Article 2*

1. Member States shall bring into force the laws, regulations and administrative provisions necessary to comply with this Directive with effect from 1 June 2008. They shall forthwith inform the Commission thereof.

When Member States adopt these measures, they shall contain a reference to this Directive or shall be accompanied by such reference on the occasion of their official publication. The methods of making such reference shall be laid down by the Member States.

2. Member States shall communicate to the Commission the text of the main provisions of national law which they adopt in the field covered by this Directive.

*Article 3*

This Directive shall enter into force on the twentieth day following its publication in the Official Journal of the European Union.

It shall apply from 1 June 2008.

Notwithstanding the second paragraph of this Article, Article 1 point 6 shall apply from 1 August 2008.

*Article 4*

This Directive is addressed to the Member States.

Done at Brussels, 18 December 2006.

*For the European Parliament*  
*The President*

*For the Council*  
*The President*

J. BORRELL FONTELLES

M. VANHANEN

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## REACH COMPLIANCE



**REGULATION (EC) No 1907/2006** of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH), establishing a European Chemicals Agency, amending Directive 1999/45/EC and repealing Council Regulation (EEC) No 793/93 and Commission Regulation (EC) No 1488/94 as well as Council Directive 76/769/EEC and Commission Directives 91/155/EEC, 93/67/EEC, 93/105/EC and 2000/21/EC.

**DIRECTIVE 2006/121/EC** of the European Parliament and of the Council of 18 December 2006 amending Council Directive 67/548/EEC on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances in order to adapt it to Regulation (EC) No 1907/2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH) and establishing a European Chemicals Agency.

### About B-Lands Consulting

**B-Lands Consulting** provides consultancy services and **LEAP AHEAD®** services designed to help worldwide organisations comply with European Union regulations.

Our company provides business friendly services on the following European Union environmental regulations:

- **REACH** (Registration, Evaluation and Authorisation of **C**hemicals) legislation
- **ELV / VHU** (End of Life Vehicle:/ Vehicules **H**ors d'**U**sage) legislation
- **EuP** (Energy Using Products) legislation
- **WEEE** (Waste **E**lectrical and **E**lectronic Equipment) legislation
- **RoHS** (Restriction of the use of **H**azardous **S**ubstances) legislation

As some EU compulsory requirements are country specific, such as for the WEEE legislation, our services are designed to adapt and conform to each country's rules and regulations, while serving both business-to-business (B2B) and business-to-consumer (B2C) companies.

We assist our customers in determining proper regulation compliance schemes, including collaborative schemes involving other businesses such as Substance Information Exchange Forums (SIEFs).

### Performed services

B-Lands Consulting offers one-stop EU regulations compliance solutions throughout all 27-EU Member States.

Services include:

- EU relevant regulations monitoring.
- Suppliers chain compliance analysis.
- Information on products labelling and other manufacturer's obligations.
- Analysis of the company products distribution system through all EU states (Subsidiaries, distributors, retailers).
- Audits on manufactured products, chemical substances & preparations (Classifications, exemption conditions, etc.).
- REACH Compliance requirements, risks assessment reports, substances full registration process
- Design of proposal for cooperation between the client and its EU distributors in any EU member state.
- Conveying products testing through a third party (Testing Labs, certifications)
- Filling of exemptions petitions on behalf of our clients to the European Commission.
- A pan European WEEE registration and recycling service for all 27-EU Member States.
- Handling of all the required paperwork. Contracts, powers of attorney are to be submitted to the client for approval and signature.

### Additional benefits are:

- The implementation and execution of smooth and on schedule compliance overhauls.
- A single and a consistent corporate solution for all EU compliance requirements.
- A single entry point for reporting to the EU bodies and registries.