

Commission Implementing Decision (EU) 2019/417 of 8 November 2018 laying down guidelines for the management of the European Union Rapid Information System ‘RAPEX’ established under Article 12 of Directive 2001/95/EC on general product safety and its notification system (notified under document C(2018) 7334)

- Article 1 The guidelines for the management of the European Union Rapid...
- Article 2 Decision 2010/15/EU is repealed.
- Article 3 This Decision is addressed to the Member States.
Signature

ANNEX

GUIDELINES FOR THE MANAGEMENT OF THE EUROPEAN UNION RAPID INFORMATION...

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PART II

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PART III

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 - 2.2. A risk assessment in three steps
 - 2.3. Some useful tips
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 - Make a sensitivity analysis of your risk assessment
 - Let others check your risk assessment
 - Document your risk assessment
 - Several hazards, several injuries — but only one risk
 - Can risks cumulate?
 - Compliance with limit values in legislation and standards
 - Specific risk assessment guidelines in specific cases
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 - Products may still present a risk even though they do...
 - 3.3. The consumer
 - 3.4. Injury scenario: Steps leading to injury(ies)

- 3.5. Severity of injury
- 3.6. Probability of injury
- 3.7. Determination of risk
4. From risk to action: how to manage risk responsibly
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 5. Determine the probability of the injury scenario.
 6. Determine the risk level.
 7. Check whether the risk level is plausible.
 8. Develop several injury scenarios to identify the highest risk of...
 9. Document and pass on your risk assessment.
6. Examples
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 - 6.3. Sensitivity analysis

Table 3 Severity of injury

Introduction

Glossary of terms

Changes to legislation: *There are currently no known outstanding effects for the Commission Implementing Decision (EU) 2019/417. (See end of Document for details)*

- (1) [OJ L 11, 15.1.2002, p. 4.](#)
- (2) [OJ L 218, 13.8.2008, p. 30.](#)
- (3) Commission Decision 2010/15/EU of 16 December 2009 laying down guidelines for the management of the Community Rapid Information System ‘RAPEX’ established under Article 12 and of the notification procedure established under Article 11 of Directive 2001/95/EC (the General Product Safety Directive) ([OJ L 22, 26.1.2010, p. 1.](#))

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There are currently no known outstanding effects for the Commission Implementing Decision (EU) 2019/417.