Commission Regulation (EC) No 627/2006 of 21 April 2006 implementing Regulation (EC) No 2065/2003 of the European Parliament and of the Council as regards quality criteria for validated analytical methods for sampling, identification and characterisation of primary smoke products

COMMISSION REGULATION (EC) No 627/2006

of 21 April 2006

implementing Regulation (EC) No 2065/2003 of the European Parliament and of the Council as regards quality criteria for validated analytical methods for sampling, identification and characterisation of primary smoke products

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 2065/2003 of the European Parliament and the Council of 10 November 2003 on smoke flavourings used or intended for use in or on foods[(1)] and in particular Article 17(3) thereof,

Whereas:

(1) Regulation (EC) No 2065/2003 lays down provisions for the establishment of a list of primary products authorised for use as such in or on foods and for the production of smoke flavourings for use in or on foods within the Community. This list shall, among other things, contain a clear description and characterisation of each primary product.

(2) Detailed information about the qualitative and quantitative chemical composition of the primary product is necessary for the scientific evaluation. The portions which have not been identified, i.e. the amount of substances whose chemical structure is not known, should be as small as possible.

(3) It is therefore necessary to establish minimum performance criteria, in this context referred to as quality criteria, to which the method of analysis shall comply in order to ensure that laboratories use methods with the necessary level of performance.

(4) Smoked foods in general give rise to health concerns, especially with respect to the possible presence of polycyclic aromatic hydrocarbons (PAHs).

(5) The person who intends to place primary products on the market should submit all the information necessary for the safety assessment. This information should contain a proposed validated method for sampling, identification and characterisation of the primary product.

(6) Regulation (EC) No 882/2004 of the European Parliament and the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules[(2)], lays down general requirements for methods of sampling and analysis.

Changes to legislation: *There are outstanding changes not yet made to Commission Regulation (EC) No 627/2006. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes*

(7) The Scientific Committee on Food (SCF) named 15 PAHs as potentially genotoxic and carcinogenic to humans in an opinion on the risks to human health of PAH in food, expressed on 4 December 2002[3]. They represent a priority group in the assessment of the risk of long-term adverse health effects following dietary intake of PAHs. Their presence in primary products should — as a consequence — be analysed.

(8) The Institute for Reference Materials and Measurements (IRMM) of the Commission Directorate General Joint Research Centre carried out collaborative studies for analysing the chemical composition of primary products and for quantifying the concentration of the 15 PAHs therein. The results of these trials are in part published in the Report on the Collaborative Trial for Validation of two Methods for the Quantification of Polycyclic Aromatic Hydrocarbons in Primary Smoke Condensates[4].

(9) To describe the precision of the method the repeatability standard deviation as defined in ISO 5725-1[5] is required. It should be estimated using data from a single-laboratory validation exercise giving S_i as described in the Harmonized Guidelines for Single-Laboratory Validation of Methods of Analysis[6] or a collaborative trial giving S_r and S_R as described in the Protocol for the design, conduct and interpretation of method-performance studies[7].

(10) A full validation of methods to analyse the composition of primary products, with a maximum of compounds identified, is not achievable. The high number of analytes gives rise to incalculable amount of work which is impractical. If however mass spectrometry is used for the detection of compounds, the resulting mass spectra can be compared to published data[8] or to mass spectral libraries and a tentative identification of the compounds can be achieved.

(11) Based on the results obtained in the inter-laboratory validation study on PAHs and following Commission Decision 2002/657/EC[9], minimum quality criteria for any suitable analytical method for determination of PAHs in all primary products have been proposed.

(12) Following the recommendation given in the ISO, IUPAC, and AOAC International Harmonized Guidelines for the Use of Recovery Information in Analytical Measurement, the analytical results should be corrected for recovery.

(13) The European Food Safety Authority has given scientific and technical assistance for the elaboration of the quality criteria for validated methods for identification and characterisation of primary smoke products laid down in this regulation.

(14) The quality criteria can be adapted, to take into account advances in scientific and technological knowledge.

(15) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

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Article 1

The quality criteria for validated analytical methods for sampling, identification and characterisation of primary smoke products, as referred to in point 4 of Annex II to Regulation (EC) No 2065/2003, shall be as set out in the Annex to this Regulation.

Article 2

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

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 21 April 2006.

For the Commission

Markos KYPRIANOU

Member of the Commission

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ANNEX

Quality criteria for validated analytical methods for sampling, identification and characterisation of primary smoke products

1. Sampling

The basic requirement is to obtain a representative and homogeneous laboratory sample.

The analyst shall ensure that samples do not become contaminated during sample preparation. Containers have to be rinsed with high purity acetone or hexane (p.A., HLPC grade or equivalent) before use to minimise the risk of contamination. Wherever possible, apparatus coming into contact with the sample shall be made of inert materials e.g. glass or polished stainless steel. Plastics such as polypropylene are to be avoided, because the analyte can adsorb onto these materials.

All of the sample material received by the laboratory is to be used for the preparation of test material. Only very finely homogenised samples give reproducible results.

There are many satisfactory specific sample preparation procedures which may be used.

2. Identification and characterisation

2.1. Definitions

For the purpose of this Annex the following definitions shall apply:

Solvent free mass	:	The mass of the material after abstraction of the solvent, which normally is water.
Volatile fraction	:	The part of the solvent free mass, which is volatile and analysable by gas chromatography.
Identification of a primary product	:	Results of a descriptive analysis, which identifies substances present in the primary product.
Characterisation of a primary product	:	Identification of the major physical-chemical fractions and quantification and identification of the chemical constituents.
LOQ	:	Limit of Quantification
LOD	:	Limit of Detection
S_i	:	The single-laboratory standard deviation, calculated from results generated under repeatability conditions as defined in ISO standard 5725-1[(10)] (= repeatability standard deviation estimated in a single laboratory approach according to the Harmonized Guidelines for Single-Laboratory Validation of Methods of Analysis[(11)]).
S_r	:	The average within laboratory standard deviation, calculated from results generated under repeatability conditions as defined in ISO standard 5725-1[(10)] in a collaborative trial with a minimum of eight laboratories conducted according to the Protocol for the Design, Conduct and Interpretation of Method-Performance Studies[(12)].
S_R	:	The between laboratory standard deviation, calculated from results under reproducibility conditions as defined in ISO standard 5725-1[(10)] and according to the Protocol for the Design, Conduct and Interpretation of Method-Performance Studies[(12)].
RSD_i	:	Relative single-laboratory repeatability standard deviation (S_i expressed in percent of the measured value),