Testing and Validation of Polycyclic Aromatic Hydrocarbons (PAH) in the course of GS-Mark Certification

1 Aim

Products (technical work equipment and consumer products) must comply with legal requirements to avoid health risks, e.g. §§ 30 & 31 of the LFBG, the Prohibition of Chemicals Act, and the § 4 of the GPSG. With this document and the attached test procedure the requirement regarding PAH in products is substantiated. In addition, the test method is harmonized for the validation by GS bodies.

The Board for Technical Work Equipment and Consumer Products (AtAV) on Nov. 20, 2007 has demanded that the consideration of the test for PAH for the GS certification of products has to be mandatory. The implementation takes place by means of this PAH document.

2 Basics

The main reasons for PAH contamination in materials is the use of:
- PAH contaminated softening oils in rubber and flexible (soft) plastics
- PAH contaminated soot as a black pigment dye in rubber and plastics

PAH contamination could be proven so far not only in rubber, but also in various plastic materials, such as ABS, PP and various lacquer / coating as well as in natural materials.

3 Procedure

The following steps are to be considered by the GS body at new GS mark awards as well as in the frame of monitoring of existing GS mark certificates:

1. Risk analysis
2. Categorization
3. Test and evaluation

3.1 Risk Analysis

The GS body has to perform a risk analysis and at the same time has to stipulate which (grip) surfaces are possible for a test and which are not as well as to document this. This means, that the GS body first of all has to stipulate for which (grip) surfaces the requirements of the PAH document apply (stipulation of the PAH relevance). As the case may be, the risk analysis can be omitted, if the circle of exchange of experience has already determined a procedure for the product and the product group respectively regarding the (grip) surfaces to be tested. The reference to the EK regulation is to be documented accordingly.

Materials which are not accessible or only accessible with the help of tools needn’t to be tested.
Basically all (grip) surfaces are to be considered, which during intended use or foreseeable misapplication (however not misuse) can be touched or put into the mouth\(^1\)

Materials which may contain PAH are e.g. elastomers (plastics and rubber materials) as well as black or dark coloured polymers.

### 3.2 Categorization

Depending on the result of the risk analysis afterwards the respective products are to be categorized (see table 1) and to be tested regarding the actual content of PAH according to the method of analysis mentioned below. Existing rest reports can be considered provided that the ZEK Principle Decision ZEK-GB-2000-01 and the requirements of this PAH document are observed. The categorization can be omitted, if the respective circle of exchange of experiences for a product and a product group respectively has already stipulated a categorization of the (grip) surfaces. Definitions to products and/or product groups of the individual exchange of experience circles are published on the ZLS homepage and to be valid starting from the time of the publication.

The limits for PAH that must be met for materials of consumer goods are stated in table 1.

The regulations of this document regarding the PAH content do not apply if in other laws corresponding or advanced requirements to the PAH content are given. This applies only for the material or component and assembly group respectively and not for the whole product. Materials and parts of the product which are not covered by other laws are to be evaluated in the frame of the GS mark award according to the specifications of the PAH document\(^2\).

It must be ensured that the detection limit of 0.2 mg/kg for each of the PAH single component can actually be proved by the test procedure.

In the sum of all 16 PAH according to EPA only those PAH components are take into account, which have been specified in the material over the 0.2 mg/kg.

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\(^1\) In order to guarantee a consistent and adequate procedure for the GS mark award not all free accessible surface should be checked generally. Intention of this document is that actually only relevant (grip) surfaces are considered. It is no target-oriented to test all product parts or surfaces “for safety”.

\(^2\) Example: Water-conducting in coffee machines which are in contact with food (e.g. water etc.) are subjected to the law LFGB, and therefore excluded from the practice of the PAH document. Grip surfaces of the coffee machine however have to be evaluated according to the requirements of the PAH document further on.
Table 1: Limits for PAH content for material of (grip) surfaces, which are to be categorized on account of the results of the risk analysis.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Category 1</th>
<th>Category 2</th>
<th>Category 3</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Materials intended to be put in the mouth or materials of toys for children aged &lt; 36 months with intended skin contact 2)</td>
<td>Materials not covered by category 1, with foreseeable skin contact for longer than 30 seconds (long-term skin contact)</td>
<td>Materials not covered by category 1 or 2 with foreseeable skin contact up to 30 seconds (short term skin contact)</td>
</tr>
<tr>
<td>Benzo[a]pyrene mg/kg</td>
<td>Not detectable (&lt; 0.2) 1)</td>
<td>1</td>
<td>20</td>
</tr>
<tr>
<td>Sum 16 PAH (EPA) mg/kg</td>
<td>Not detectable (&lt; 0.2) 1)</td>
<td>10</td>
<td>200</td>
</tr>
</tbody>
</table>

1) If the limits of category 1 are surpassed but the limits of category 2 still met, the confirmation of suitability of contact with foodstuff or the oral mucosa can be verified by an additional specific migration test of the PAH components according to EN 1186, ff. and § 64 LFBG 80.30-1. The results of the migration test shall be evaluated according to law criteria for foodstuff.

2) according to the regulations of the EK2.

3.3 Test and Evaluation

The steps of the preparation of sampling, extraction of the PAH, purification of the extract, identification and quantitation are described in the attached test instruction and have to be applied by all performing test laboratories in equal measure.

The GS body evaluates the test result and decides whether the GS mark can be awarded subject to the other requirements.

4 Transition Regulations

4.1 Transition Periods

At the award of the GS mark for products this document is mandatory from 1st April, 2008 onwards (based on the stipulation of the AtAV meeting dated 2007-11-20.

As the test regarding the PAH contents in products represents a comprehensive requirement to nearly all members of the circles of exchange of experience the following procedure is determined by the ZLS:

4.1.1. GS mark certificates, date of issue from 2008-04-01 onwards (including ongoing projects, which will be completed after 2008-04-01)
(This document supersedes the document ZEK 01.1-08 with the date of publications)

Mandatory application on the PAH document ZEK 01-08 from 1st April, 2008 onwards and ZEK 01.1-08 respectively from date of publication onwards (exception: see 4.1.3).

4.1.2. GS mark certificates, issued before 1st April, 2008

For the time being, existing GS mark certificates remain valid.

Within the framework of regular control measures for monitoring the production (at the latest within 1 year and in those cases respectively, when the regular control period is two years, within 2 years) the requirements of paragraph 3 of the EK document according to the risk analysis have to be considered, regardless whether the product was inspected in the factory or not. In case it can be concluded that the respective requirements are not met, the certificate has to be withdrawn immediately The ZEK Principle Decision ZEK-GB-2006-01 is to be observed.

4.1.3 New issue of existing GS mark certificates – Exceptional Rules

An immediate consideration is not necessary in case of a new issue of an existing GS mark certificate for the following reasons

In case that the trade name is changed normally new GS mark certificates are issued. As the product is not modified regarding the construction as well as other properties and the new issue of the GS mark certificate is more or less a mere formality, a consideration of the requirements of the PAH resolution is only necessary when the control measure for monitoring the production takes place.

(Note: With the new issue of the GS mark certificate the already specified periods for the control measures of the manufacture of the product do not change.)

The same applies in an analogous manner at a relocation of the GS mark certificate owner provided that the product is not changed concerning all properties and no additional safety-related examination of the product is necessary.

For the area of the secondary certificates (OEM certificates) the same procedure mentioned above can be applied. In these cases an examination according to the specifications of the PAH document until the next control measure for the manufacture of the product is mandatory. This has to take place according the already determined periods of the “main certificate” and until 31st March, 2009 at the latest. Control periods in excess thereof are not allowed in these cases.

Concerning the OEM certificates and therefore also for the „main certificates“ the PAH document has to be applied – as stipulated – until 31st March, 2009 at the latest.
Test Instruction
Harmonised Method for Determination of Polycyclic Aromatic Hydrocarbons (PAH) in Plastic Sampling

1 Aim
To identify PAH Poly cyclic Aromatic Hydrocarbons in plastic sampling (material).

2.1 Short Description of the Procedure

2.1.1 General Procedure
A representative partial sample is taken from the material and cut with scissors, side cutters etc. into particles of max 2-3 mm in size. Of that 500 mg shall be weighed for extraction. Use 20 ml toluene mixed with internal standard. Place for 1 hour into the ultrasonic bath at a temperature of 60° C for extraction.

After cooling to room temperature take an aliquot from the extract. At plastic and rubber material respectively at which matrix problems occur during the examination, an additional column-type chromatographic purification step is performed.

The quantification is done with a gas chromatograph with a mass specific detector (GC-MSD) in SIM method.

2.1.1 Procedure with Smaller Quantities
If the total mass of the material which can be examined should fall below 500 mg, the following is valid:

Identical materials of the product can be united and regarded as a sample.

Additional product samples may not be used however.

If less than 50 mg material is available for individual samples, these are not examined.

If the available mass of the cut up material amounts to only between 50 mg and 500 mg, then the sample is to be examined after 2.1.1, and the Toluol quantity is to be converted and/or adapted proportionally. The actual mass of the sample is to be specified accordingly in the test report.

2.2 Equipment

- Ultrasonic bath: minimum power is 200 W for a bath surface of 706 cm² which compares to 0.28 W/cm² without basket with internal or external thermostat (temperature controller)
- Gas chromatograph with mass selective detector

Date: 2008-06-23
2.3 Chemicals and Solutions

2.3.1 Chemicals

- Toluene
- Internal standards: Standard 1: naphthalene - d8
  Standard 2: pyren-d10 or anthracene-d10, or phenanthrene-d10
  Standard 3: benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene

At least 3 internal standards shall be used to be mixed with toluene as extraction agent.

- External standard: 16 PAH-substances according to EPA, mixed or individual.
- Petroleum ether
- Silicia gel
- Sodium sulphate

2.3.2 Calibration Solutions

The concentration of the calibration solutions must be selected in such way that a 3-point-calibration covers the range of the samples from 0.1 to 10 mg/kg.

This compares to a concentration range in the calibration solutions of 2.5 to 250 ng/ml.

3.1 Sample Preparation

Take a representative partial sample from the material. The dimensions of the particles shall not be larger than 2-3 mm. Use scissors, side cutters, pliers to generate the particles from the sample.

3.1.1 Extraction

500 mg of the sample shall be mixed with the toluene (already mixed with the internal standards) in a (flange-)glass. This is then placed into the ultrasonic bath for 1 hour at constant 60°C for extraction. Place the glasses into the bath standing without the basket or hang them. Afterwards the glasses are taken out and an aliquot of the extract is taken after cooling to room temperature and short shaking. Measure directly from the aliquot or after thinning with toluene.

3.1.2 Column-type chromatographic extract purification step

At some plastic and rubber products respectively, especially such, which are dissolved with toluene to a very large degree under the described extraction conditions, a purification of the extract with silicia gel adsorption chromatography is necessary.

Hereto a clean-up column with “Hahnschliff” (ca. 220 x 15 mm) with fibreglass, 4 g silica gel and 1 cm sodium sulphate. The silicia gel first is to be deactivated by addition of 10 % water (in the glass flask the silicia gel is mixed with the appropriate amount of water and afterwards 1 h homogenized on the rotary vaporizer at 760 Torr and room temperature. The silicia gel then can be stored in the closed flask at room temperature).
The conditioning of the packed column is done with 10 ml petroleum ether. Then the toluene extract aliquot in the rotary vaporizer is reduced to ca. 1 ml and given upon the column. The pointed piston is rinsed with ca. 20 ml elution agent which is also transferred to the clean-up column. The elution is done with 50 ml petroleum ether. The collected petroleum ether eluat is spiked with 1 ml toluene and reduced at the TurboVap with nitrogen to ca. 1 ml. Afterwards it is replenished with toluene to a defined volume and the extract then is analysed by means of GC-MS.

3.2 Measurement Procedure

The determination method to be used is the gas chromatography with mass selective detection in SIM Mode. The following 16 PAH must be identified according to EPA:

- NAPHTHALENE
- ACENAPHTHYLENE
- ACENAPHTHENE
- FLUORENE
- PHENANTHRENE
- ANTHRACENE
- FLUORANTHENE
- PYRENE
- BENZO(a)ANTHRACENE
- CHRYLENE
- INDENO(1,2,3-cd)PYRENE
- BENZO(b)FLUORANTHENE
- BENZO(k)FLUORANTHENE
- BENZO(a)PYRENE
- DIBENZO(a,h)ANTHRACENE
- BENZO(g,h,i)PERYLENE

3.2.1 Gas Chromatographic Measurement Conditions

The equipment parameters (temperature, columns, mass traces) are to be decided by the lab respectively, by its staff analyts.

3.2.2 Evaluation

- Internal standards: use of at least 3 internal standards. They and the correction range are defined as follows:

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Internal standards with recommended reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>NAPHTHALENE</td>
<td>naphtalene - d8</td>
</tr>
<tr>
<td>ACENAPHTHYLENE</td>
<td>pyrene-d10 or anthracene-d10, or phenanthrene-d10</td>
</tr>
<tr>
<td>ACENAPHTHENE</td>
<td>pyrene-d10 or anthracene-d10, or phenanthrene-d10</td>
</tr>
<tr>
<td>FLUORENE</td>
<td>pyrene-d10 or anthracene-d0, or phenanthrene-dl 0</td>
</tr>
<tr>
<td>PHENANTHRENE</td>
<td>pyrene-d10 or anthracene-d10, or phenanthrene-d10</td>
</tr>
</tbody>
</table>
d10 ANTHRACENE pyrene-d10 or anthracene-d10, or phenanthrene-d10
FLUORANTHENE pyrene-d10 or anthracene-d10, or phenanthrene-d10
PYRENE pyrene-d10 or anthracene-d10, or phenanthrene-d10
BENZO(a)ANTHRACENE pyrene-d10 or anthracene-d10, or phenanthrene-d10
CHRYSENE pyrene-d10 or anthracene-d10, or phenanthrene-d10
INDENO(l,2,3-cd)PYRENE benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene
BENZO(b)FLUORANTHENE benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene
BENZO(k)FLUORANTHENE benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene
BENZO(a)PYRENE benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene
INDENO(1,2,3-cd)PyRENE benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene
DIBENZO(a,h)ANTHRACENE benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene
BENZO(g,h,i)PERYLENE benzo(a)pyrene-d12 or perylene-d12 or triphenylbenzene

* External calibration: for each single PAH at least one 3-point-calibration with reference to the afore mentioned internal standardization must be done. Hereby a work range of 0.1 - 10 mg/kg is recommended.

* Concentrations above the calibration range can be determined by thinning of the extract.

3.2.3 Determination limit
The determination limit for material samples is 0.2 mg/kg per parameter.

3.3 Special characteristics
Based on its relative volatility against the other 15 PAH (according to EPA), naphthalene represents a parameter difficult to evaluate in close to skin products.
Experience of the testing bodies show that loss of naphthalene as well as secondary contamination can be found.
The developed naphthalene result will always only show the momentary situation of the test sample at the time of measurement.

Attachment: Gas Chromatic Measurement Conditions (informative)
Injected amount: 1|j|, pulsed splitlos
Column: HT8 25m, ID 0,22mm, film thickness 0,25um
Injector temperature: 280°C
Transfer line temperature: 260°C
Initial temperature: 5CTC
Initial time: 2min
Heating rate: 11°C/min
Final temperature: 320°C
Final time: 8 min
<table>
<thead>
<tr>
<th>Parameter</th>
<th>RT ca min</th>
<th>MOL-MASS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Naphthalene</td>
<td>10,5</td>
<td>128</td>
</tr>
<tr>
<td>Acenaphthylene</td>
<td>14,5</td>
<td>152</td>
</tr>
<tr>
<td>Acenaphthene</td>
<td>14,8</td>
<td>153</td>
</tr>
<tr>
<td>Fluorene</td>
<td>16,0</td>
<td>166</td>
</tr>
<tr>
<td>Phenanthrene</td>
<td>18,3</td>
<td>178</td>
</tr>
<tr>
<td>Anthracene</td>
<td>18,4</td>
<td>178</td>
</tr>
<tr>
<td>Fluoranthene</td>
<td>21,1</td>
<td>202</td>
</tr>
<tr>
<td>Pyrene</td>
<td>21,7</td>
<td>202</td>
</tr>
<tr>
<td>Chrysene</td>
<td>24,5</td>
<td>228</td>
</tr>
<tr>
<td>Benzo(a)anthracene</td>
<td>24,6</td>
<td>228</td>
</tr>
<tr>
<td>Benzo(b)fluoranthene</td>
<td>27,0</td>
<td>252</td>
</tr>
<tr>
<td>B9nzo(k)fluoranthene</td>
<td>27,0</td>
<td>252</td>
</tr>
<tr>
<td>Benzo(a)pyrene</td>
<td>27,8</td>
<td>252</td>
</tr>
<tr>
<td>Indeno(1-2,3-cd)pyrene</td>
<td>31,2</td>
<td>276</td>
</tr>
<tr>
<td>Dibenzo(ah)anthracene</td>
<td>31,5</td>
<td>278</td>
</tr>
<tr>
<td>Benzo(ghi)perylenene</td>
<td>32,7</td>
<td>276</td>
</tr>
</tbody>
</table>