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EFSA Panel on Food Contact Materials, Enzymes, Flavourings and Processing Aids (CEF), Vittorio Silano, Claudia Bolognesi, Laurence Castle, Kevin Chipman, Jean-Pierre Cravedi, Karl-Heinz Engel, Paul Fowler, Roland Franz, Konrad Grob, Rainer Gürtler, Trine Husøy, Sirpa Kärenlampi, Maria Rosaria Milana, Karla Pfaff, Gilles Riviere, Jannavi Srinivasan, Maria de Fátima Tavares Poças, Christina Tlustos, Detlef Wölfle, Holger Zorn, Romualdo Benigni, Mona-Lise Binderup, Leon Brimer, Francesca Marcon, Daniel Marzin, Pasquale Mosesso, Gerard Mulder, Agneta Oskarsson, Camilla Svendsen, Jan van Benthem, Maria Anastassiadou, Maria Carfi and Wim Mennes

Abstract

The Panel on Food Contact Materials, Enzymes, Flavourings and Processing Aids of the EFSA was requested to consider evaluations of flavouring substances assessed since 2000 by the Joint FAO/WHO Expert Committee on Food Additives (JECFA) and to decide whether further evaluation is necessary, as laid down in Commission Regulation (EC) No 1565/2000. The present consideration concerns a group of 22 pyridine, pyrrole and quinoline derivatives evaluated by JECFA (63rd meeting). The revision of this consideration is made since additional genotoxicity data have become available for 6-methylquinoline [FL-no: 14.042]. The genotoxicity data available rule out the concern with respect to genotoxicity and accordingly the substance is evaluated through the Procedure. For all 22 substances [FL-no: 13.134, 14.001, 14.004, 14.007, 14.030, 14.038, 14.039, 14.041, 14.042, 14.045, 14.046, 14.047, 14.058, 14.059, 14.060, 14.061, 14.065, 14.066, 14.068, 14.071, 14.072 and 14.164] considered in this Flavouring Group Evaluation (FGE), the Panel agrees with the JECFA conclusion, 'No safety concern at estimated levels of intake as flavouring substances' based on the Maxiimised Survey-derived Daily Intake (MSDI) approach. Besides the safety assessment of these flavouring substances, the specifications for the materials of commerce have also been evaluated, and the information is considered adequate for all the substances. For the following substances [FL-no: 13.134, 14.001, 14.030, 14.041, 14.042, 14.058, 14.072], the Industry has submitted use levels for normal and maximum use. For the remaining 15 substances, use levels are needed to calculate the modified Theoretical Added Maximum Daily Intakes (mTAMDis) in order to identify those flavouring substances that need more refined exposure assessment and to finalise the evaluation.

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Keywords: pyridine, FGE.77, pyrrole, quinoline, JECFA, 63rd meeting, FGE.24Rev2

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1. Introduction

1.1. Background and Terms of Reference as provided by the requestor

1.1.1. Background

The use of flavourings is regulated under Regulation (EC) No 1334/2008 of the European Parliament and Council of 16 December 2008 on flavourings and certain food ingredients with flavouring properties for use in and on foods. On the basis of Article 9(a) of this Regulation, an evaluation and approval are required for flavouring substances.

The Union list of flavourings and source materials was established by Commission Implementing Regulation (EC) No 872/2012. The list contains flavouring substances for which the scientific evaluation should be completed in accordance with Commission Regulation (EC) No 1565/2000.

The substance 6-methylquinoline [FL-no: 14.042] is one of these substances. New genotoxicity data on it was evaluated in the EFSA opinion FGE.77Rev1 (January 2014) but EFSA concluded that this new data did not clear the concern with respect to genotoxicity in vitro and indicated that studies in vivo would be necessary to address the potential for genotoxicity. EFSA evaluated FGE.77 again in January 2015 (FGE.77Rev2) due to new studies on other substances of this group, although at that time no new data on FL-no: 14.042 was available.

On 27 January 2016 the Industry submitted a new dossier with the requested studies by EFSA.

1.1.2. Terms of Reference

The European Commission requests the European Food Safety Authority (EFSA) to evaluate this new information and, depending on the outcome, proceed to the full evaluation of this flavouring substance in accordance with Commission Regulation (EC) No 1565/2000.

2. Assessment

The approach used by EFSA for safety evaluation of flavouring substances is referred to in Commission Regulation (EC) No 1565/2000, hereafter named the ‘EFSA Procedure’. This Procedure is based on the opinion of the Scientific Committee on Food (SCF) (SCF, 1999), which has been derived from the evaluation procedure developed by the Joint FAO/WHO Expert Committee on Food Additives (JECFA, 1995, 1996, 1997, 1999), hereafter named the ‘JECFA Procedure’. The Panel on Food Contact Materials, Enzymes, Flavourings and Processing Aids (the Panel) compares the JECFA evaluation of structurally related substances with the result of a corresponding EFSA evaluation, focussing on specifications, intake estimations and toxicity data, especially genotoxicity data. The evaluations by EFSA will conclude whether the flavouring substances are of no safety concern at their estimated levels of intake, whether additional data are required or whether certain substances should not be evaluated through the EFSA Procedure.

The following issues are of special importance.

Intake

In its evaluation, the Panel as a default uses the Maximised Survey-derived Daily Intake (MSDI) approach to estimate the per capita intakes of the flavouring substances in Europe.

In its evaluation, JECFA includes intake estimates based on the MSDI approach derived from both European and USA production figures. The higher of the two MSDI figures is used in the evaluation by JECFA. It is noted that in several cases, only the MSDI figures from the USA were available, meaning that certain flavouring substances have been evaluated by JECFA only on the basis of these figures. For Register substances for which this is the case the Panel will need European Union (EU) production figures in order to finalise the evaluation.

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1 Regulation (EC) No 1334/2008 of the European Parliament and of the Council of 16 December 2008 on flavourings and certain food ingredients with flavouring properties for use in and on foods and amending Council Regulation (EEC) No 1601/91, Regulations (EC) No 2232/96 and (EC) No 110/2008 and Directive 2000/13/EC. OJ L 354, 31.12.2008, p. 34–50.

2 Commission implementing Regulation (EU) No 872/2012 of 1 October 2012 adopting the list of flavouring substances provided for by Regulation (EC) No 2232/96 of the European Parliament and of the Council, introducing it in Annex I to Regulation (EC) No 1334/2008 of the European Parliament and of the Council and repealing Commission Regulation (EC) No 1565/2000 and Commission Decision 1999/217/EC. OJ L 267, 2.10.2012, p. 1–161.

3 Commission Regulation No 1565/2000 of 18 July 2000 laying down the measures necessary for the adoption of an evaluation programme in application of Regulation (EC) No 2232/96. OJ L 180, 19.7.2000, p. 8–16.
When the Panel examined the information provided by the European Flavour Industry on the use levels in various foods, it appeared obvious that the MSDI approach in a number of cases would grossly underestimate the intake by regular consumers of products flavoured at the use level reported by the Industry, especially in those cases where the annual production values were reported to be small. In consequence, the Panel had reservations about the data on use and use levels provided and the intake estimates obtained by the MSDI approach. It is noted that JECFA, at its 65th meeting considered ‘how to improve the identification and assessment of flavouring agents, for which the MSDI estimates may be substantially lower than the dietary exposures that would be estimated from the anticipated average use levels in foods’ (JECFA, 2006b).

In the absence of more accurate information that would enable the Panel to make a more realistic estimate of the intakes of the flavouring substances, the Panel has decided also to perform an estimate of the daily intakes per person using a modified Theoretical Added Maximum Daily Intake (mTAMDI) approach based on the normal use levels reported by Industry.

As information on use levels for the flavouring substances has not been requested by JECFA or has not otherwise been provided to the Panel, it is not possible to estimate the daily intakes using the mTAMDI approach for the substances evaluated by JECFA. The Panel will need information on use levels in order to finalise the evaluation.

**Threshold of 1.5 μg/person per day (Step B5) used by JECFA**

JECFA uses the threshold of concern of 1.5 μg/person per day as part of the evaluation procedure:

‘The Committee noted that this value was based on a risk analysis of known carcinogens which involved several conservative assumptions. The use of this value was supported by additional information on developmental toxicity, neurotoxicity and immunotoxicity. In the judgement of the Committee, flavouring substances for which insufficient data are available for them to be evaluated using earlier steps in the Procedure, but for which the intake would not exceed 1.5 μg per person per day would not be expected to present a safety concern. The Committee recommended that the Procedure for the Safety Evaluation of Flavouring Agents used at the forty-sixth meeting be amended to include the last step on the right-hand side of the original procedure (“Do the condition of use result in an intake greater than 1.5 μg per day?”) (JECFA, 1999).

In line with the Opinion expressed by the SCF (SCF, 1999), the Panel does not make use of this threshold of 1.5 μg/person per day.

**Genotoxicity**

As reflected in the Opinion of SCF (SCF, 1999), the Panel has in its evaluation focussed on a possible genotoxic potential of the flavouring substances or of structurally related substances. Generally, substances for which the Panel has concluded that there is an indication of genotoxic potential in vitro, will not be evaluated using the EFSA Procedure until further genotoxicity data are provided. Substances for which a genotoxic potential in vivo has been concluded, will not be evaluated through the Procedure.

**Specifications**

Regarding specifications, the evaluation by the Panel could lead to a different opinion than that of JECFA.

**Structural relationship**

In the consideration of the JECFA evaluated substances, the Panel will examine the structural relationship and metabolism features of the substances within the flavouring group and compare this with the corresponding Flavouring Group Evaluation (FGE).

**2.1. History of the evaluation of the substances in the present FGE**

JECFA has evaluated a group of 22 flavouring substances consisting of pyridine, pyrrole and quinoline derivatives (JECFA, 2006a).

These 22 substances were considered by EFSA in FGE.77, in which the Panel concluded that additional toxicity data were needed for seven substances [FL-no: 13.134, 14.001, 14.041, 14.045, 14.046, 14.047 and 14.068] as no adequate toxicity studies were available from which a no observed adverse effect level (NOAEL) could be established, neither on the substances nor on supporting substances. The Panel also concluded, contrary to JECFA, that 6-methylquinoline [FL-no: 14.042] should not be evaluated through the Procedure due to concern with respect to genotoxicity in vitro.
In the first Revision of FGE.77, FGE.77Rev1, additional toxicity data were provided for isoquinoline [FL-no: 14.001], pyrrole [FL-no: 14.041] and 2-acetylpyrrole [FL-no: 14.047]; the toxicity data on 2-acetylpyrrole also cover 2-propionylpyrrole [FL-no: 14.068]. The main studies provided were for each substance a 90-day study. Furthermore, additional genotoxicity data for 6-methylquinoline [FL-no: 14.042] became available. EU production volumes were provided for four substances, [FL-no: 14.045, 14.058, 14.059 and 14.164] for which the evaluation could not be finalised previously, due to lack of these data. Based on these newly submitted EU production volumes (IOFI, 2013), the substances were already evaluated in FGE.964 (EFSA CEF Panel, 2011), but for the sake of completion, the information was included here as well. Finally, information on solubility was provided for six substances [FL-no: 13.134, 14.007, 14.030, 14.038, 14.045 and 14.046] since the previous evaluation of FGE.77.

The second revision of FGE.77, FGE.77Rev2, included additional toxicity data, among which an oral 90-day study provided for 1-furfurylpyrrole [FL-no: 13.134]. The data were intended to cover the re-evaluation of this substance as well as 2-acetyl-1-ethylpyrrole [FL-no: 14.045] and 2-acetyl-1-methylpyrrole [FL-no: 14.046]. A search in open literature was conducted for metabolism, genotoxicity and toxicity for 1-furfurylpyrrole. This search did not reveal any pertinent new information on the substance.

The present revision of FGE.77, FGE.77Rev3, includes additional genotoxicity and toxicity data for 6-methylquinoline [FL-no: 14.042].

3. Presentation of the substances in the JECFA flavouring group

3.1. Description

3.1.1. JECFA status

At the 63rd meeting, JECFA has evaluated a group of 22 flavouring substances consisting of pyridine, pyrrole and quinoline derivatives (JECFA, 2005b, 2006a).

3.1.2. EFSA considerations

The Panel concluded that all the substances in the JECFA flavouring group of pyridine, pyrrole and quinoline derivatives are structurally related to the group of pyridine, pyrrole, indole and quinoline derivatives from chemical group 28 evaluated by EFSA in the Flavouring Group Evaluation 24, Revision 2 (FGE.24Rev2) (EFSA CEF Panel, 2013).

3.2. Isomers

None of the 22 flavouring substances in the group of pyridine, pyrrole and quinoline derivatives has possibility for stereoisomerism.

3.3. Specifications

3.3.1. Status

The JECFA specifications are available for all 22 substances (JECFA, 2005a) (see Table 1).

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4 Consideration of 88 flavouring substances considered by EFSA for which EU production volumes/anticipated production volumes have been submitted on request by DG SANCO.
3.3.2. EFSA considerations

The specifications are considered adequate for all 22 substances.

4. Assessment

4.1. Intake estimation

4.1.1. Status

For all 22 substances, evaluated through the JECFA Procedure, production volumes, based on which MDSI values can be calculated, are available for the EU (see Table D.2).

4.1.2. EFSA considerations

For seven substances [FL-no: 13.134, 14.001, 14.030, 14.041, 14.042, 14.058, 14.072], the Industry has submitted food categories\(^5\) and use levels in these for normal and maximum use (EFFA, 2012; DG SANCO, 2014) (see Table D.1, Appendix D). Based on the normal use levels, the mTAMDIs for [FL-no: 14.041] is below the threshold of concern of 1,800 µg/person per day for a structural class I substance. The mTAMDIs for [FL-no: 14.001, 14.058 and 14.072] are below the threshold of concern of 90 µg/person per day for the structural class III substances. The mTAMDIs for [FL-no: 13.134, 14.030, 14.042] are above the threshold of concern of 90 µg/person per day for the structural class III substances. For the remaining 15 substances, use levels are needed to calculate the mTAMDIs.

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\(^5\) Annex III, Commission Regulation No 1565/2000 of 18 July 2000 laying down the measures necessary for the adoption of an evaluation programme in application of Regulation (EC) No 2232/96. Official Journal of the European Communities 19.7.2000, L 180, p. 8–16.
Table 1: Summary of specification data (JECFA, 2005a)

| FL-no | JECFA-no | EU Register name                  | Structural formula | FEMA no | CoE no | CAS no | Phys.form Molecular formula | Molecular weight | Solubility(a) Solubility in ethanol(b) | Boiling point, °C(c) | Melting point, °C | Refractive Index(d) | Specific gravity(e) |
|-------|----------|-----------------------------------|--------------------|---------|--------|--------|----------------------------|------------------|----------------------------------------|----------------------|-------------------|-------------------|---------------------|
| 13.134| 1310     | 1-Furfurylpyrrole                |                    | 3284    | 2317   | 1438-94-4 | Liquid C9H9ON 147.18             | Insoluble Soluble | 76–78 (1 hPa) – NMR 98%                       |                      |                  | 1.529–1.536       | 1.078–1.084          |
| 14.001| 1303     | Isoquinoline                      |                    | 2978    | 487    | 119-65-3  | Solid C9H7N 129.16              | Slightly soluble Soluble | 242–243 27–29 NMR 97%                      |                      |                  | 1.621–1.627       | 1.097–1.103          |
| 14.004| 1304     | 3-Methylindole                    |                    | 3019    | 493    | 83-34-1   | Solid C9H7N 131.18              | Soluble Soluble | – 95–97 NMR 97%                            |                      |                  | n.a.              | n.a.               |
| 14.007| 1301     | Indole                            |                    | 2593    | 560    | 120-72-9  | Solid C9H7N 117.15              | Insoluble Soluble | n.a. 51–54 NMR 97%                         |                      |                  | n.a.              | n.a.               |
| 14.030| 1308     | 2-Pyridine methanethiol           |                    | 3232    | 2279   | 2044-73-7 | Liquid C6H7NS 125.20             | Soluble Soluble | 57–58 (0.8 hPa) – NMR 98%                    |                      |                  | 1.573–1.580       | 1.150–1.157          |
| 14.038| 1309     | 2-Acetylpymidine                  |                    | 3251    | 2315   | 1122-62-9 | Liquid C7H7ON 121.14             | Insoluble Soluble | 189–193 – IR NMR 97%                       |                      |                  | 1.518–1.524       | 1.077–1.084          |
| 14.039| 1316     | 3-Acetylpymidine                  |                    | 3424    | 2316   | 350-03-8  | Liquid C7H7ON 121.14             | Soluble Soluble | 230 – NMR 97%                             |                      |                  | 1.530–1.540       | 1.103–1.112          |
| 14.041| 1314     | Pyrrole                           |                    | 3386    | 2318   | 109-97-7  | Liquid C4H5N 67.09              | Slightly soluble Soluble | 130–131 – IR 98%                          |                      |                  | 1.507–1.510       | 0.955–0.975          |
| FL-no | FEMA no | EU Register name            | Structural formula | CAS no | Molecular formula | Molecular weight | Solubility(a) | Solubility in ethanol(b) | Boiling point, °C(c) | Melting point, °C | ID test | Assay minimum | Refractive Index(d) | Specific gravity(e) |
|-------|---------|-----------------------------|--------------------|--------|------------------|------------------|---------------|-------------------------|---------------------|-----------------|---------|---------------|-------------------|---------------------|
| 14.042| 9162-3 | 6-Methylquinoline           | ![Structure](image) | 2744   | C_{10}H_{12}N     | 143.19           | Slightly soluble| Soluble                 | 259                 | –               | NMR     | 98%           | 1.611–1.617         | 1.060–1.066          |
| 14.045| 11371  | 2-Acetyl-1-ethylpyrrole    | ![Structure](image) | 3147   | C_{8}H_{11}ON     | 137.18           | Slightly soluble| Soluble                 | 209–211              | –               | NMR     | 98%           | 1.550–1.556         | 1.052–1.058          |
| 14.046| 11373  | 2-Acetyl-1-methylpyrrole   | ![Structure](image) | 3184   | C_{8}H_{10}ON     | 123.16           | Slightly soluble| Soluble                 | 200–202              | –               | NMR     | 98%           | 1.539–1.545         | 1.037–1.043          |
| 14.047| 11721  | 2-Acetylpyrrole            | ![Structure](image) | 3202   | C_{6}H_{10}ON     | 109.13           | Soluble        | Soluble                 | n.a.                 | –               | NMR     | n.a.          | n.a.               | n.a.                |
| 14.058| 11395  | 2-Isobutylpyridine         | ![Structure](image) | 3370   | C_{9}H_{15}N      | 135.21           | Insoluble      | Soluble                 | 181                 | –               | NMR     | 97%           | 1.480–1.486         | 0.894–0.900          |
| 14.059| 11396  | 3-Isobutylpyridine         | ![Structure](image) | 3371   | C_{9}H_{15}N      | 135.21           | Insoluble      | Soluble                 | 68–68.5 (10hPa)      | –               | NMR     | 97%           | 1.488–1.494         | 0.898–0.904          |
| 14.060| 11412  | 2-Pentylpyridine           | ![Structure](image) | 3383   | C_{10}H_{15}N     | 149.24           | Insoluble      | Soluble                 | 102–107              | –               | NMR     | 97%           | 1.485–1.491         | 0.895–0.901          |
| 14.061| 11386  | 3-Ethylpyridine            | ![Structure](image) | 3394   | C_{7}H_{10}N      | 107.16           | Slightly soluble| Soluble                 | 166                 | –               | NMR     | 98%           | 1.499–1.505         | 0.951–0.957          |
| FL-no | EU Register name | Structural formula | FEMA no | CoE no | CAS no | Phys.form | Molecular formula | Molecular weight | Solubility(a) | Solubility in ethanol(b) | Boiling point, °C(c) | Melting point, °C | ID test | Assay minimum | Refractive Index(d) | Specific gravity(e) |
|-------|-----------------|---------------------|---------|--------|--------|-----------|------------------|------------------|--------------|------------------------|-------------------|----------------|--------|---------------|------------------|------------------|
| 14.065| 2,6-Dimethylpyridine | ![Formula](image) | 3540    | 11381  | 108-48-5 | Liquid   | C_7H_9N     | 107.16 | Soluble       | Soluble             | 143–145          | –               | 99%    | –             | 1.495–1.501       | 0.917–0.923       |
| 14.066| 5-Ethyl-2-methylpyridine | ![Formula](image) | 3546    | 11385  | 104-90-5 | Liquid   | C_8H_11N   | 121.18 | Slightly soluble | Soluble             | 172–175          | –               | 97%    | –             | 1.495–1.502       | 0.917–0.923       |
| 14.068| 2-Propionylpyrrole | ![Formula](image) | 3614    | 11942  | 1073-26-3 | Solid    | C_7H_9ON   | 123.16 | Slightly soluble | Soluble             | n.a.             | –               | n.a.   | –             | n.a.             | n.a.             |
| 14.071| Methyl nicotinate | ![Formula](image) | 3709    | 93-60-7 |        | Solid    | C_7H_7O_2N | 137.14 | Slightly soluble | Soluble             | n.a.             | 43–45           | –               | n.a.   | –             | n.a.             | n.a.             |
| 14.072| 2-(3-Phenylpropyl)pyridine | ![Formula](image) | 3751    | 2110-18-1 |        | Liquid   | C_14H_15N | 197.28 | Insoluble      | Soluble             | 142–143 (1 hPa) | –               | 97%    | –             | 1.558–1.563       | 1.012–1.018       |
| 14.164| 2-Propylpyridine | ![Formula](image) | 622-39-9 |        |        | Liquid   | C_8H_11N   | 121.20 | Slightly soluble | Soluble             | 169–171          | –               | 98%    | –             | 1.490–1.496       | 0.907–0.917       |

n.a.: not applicable.

(a): Solubility in water, if not otherwise stated.
(b): Solubility in 95% ethanol, if not otherwise stated.
(c): At 1013.25 hPa, if not otherwise stated.
(d): At 20°C, if not otherwise stated.
(e): At 25°C, if not otherwise stated.
4.2. Biological and toxicological data

4.2.1. Genotoxicity

4.2.1.1. Genotoxicity studies – text taken\(^6\) from JECFA (JECFA, 2006a)

**In vitro**

There was no evidence of mutagenicity in the assay for reverse mutation in bacteria when various strains of *Salmonella* Typhimurium (TA97, TA98, TA100, TA102, TA104, TA1535, TA1537, TA1538 and TM677) were incubated with indole [FL-no: 14.007] at a concentration of up to 30 \(\mu\text{mol/plate} (3,515 \mu\text{g/plate})\) (Anderson and Styles, 1978; Kaden et al., 1979; Florin et al., 1980; Ochiai et al., 1986; Vance et al., 1986; Sasagawa and Matsushima, 1991; Fujita et al., 1994), isoquinoline [FL-no: 14.001] at a concentration of up to 20,000 \(\mu\text{g/mL}\) (Sugimura et al., 1976; Nagao et al., 1977; Epler et al., 1979; Kaden et al., 1979; Sideropoulos and Specht, 1984), skatole [FL-no: 14.004] (4-methylindole) at a concentration of up to 3 \(\mu\text{mol/plate} (394 \mu\text{g/plate})\) (Florin et al., 1980; Ochiai et al., 1986; Kim et al., 1989; Sasagawa and Matsushima, 1991), pyrrole [FL-no: 14.041] at a concentration of up to 1.4 mmol/plate (93,926 \(\mu\text{g/plate})\) (Florin et al., 1980; Aeschbacher et al., 1989; Lee et al., 1994) and 3-ethylpyridine [FL-no: 14.061] at a concentration of up to 3 \(\mu\text{mol/plate} (321 \mu\text{g/plate})\) (Florin et al., 1980) with and without metabolic activation. Methyl 2-pyrrolyl ketone [FL-no: 14.047] (2-acetylpyrrole) at concentrations of 4 to 100 \(\mu\text{mol/plate}\) induced a >2-fold increase in the number of revertants/plate compared with the control when tested in *S. Typhimurium* TA98 in the absence of metabolic activation (Lee et al., 1994). However, negative results were obtained with metabolic activation as well as in *S. Typhimurium* TA100 (both with and without metabolic activation). Furthermore, no mutagenic activity was reported in either strain when incubated with methyl 2-pyrrolyl ketone at a concentration of up to 200 \(\mu\text{g/mL}\) with and without metabolic activation (Wang et al., 1994). 6-Methylquinoline [FL-no: 14.042] at a concentration of 3.3 to 3,600 \(\mu\text{g/plate}\) gave uniformly positive results in the presence of metabolic activation (Sugimura et al., 1976; Nagao et al., 1977; Dong et al., 1978; Wild et al., 1983; Takahashi et al., 1988; Debnath et al., 1992; Zeiger et al., 1992). Methylquinolines, tested at a concentration of 400 \(\mu\text{g/plate}\), showed a potent bactericidal or bacteriostatic effect, with only 6% survival of *S. Typhimurium* TA100 treated with 6-methylquinoline (Dong et al., 1978).

There was no evidence of mutagenicity when *Escherichia coli* (strains WP2 uvrA/pKM101, SD-4-73, or B/r HCR\(^+\)) were incubated with indole [FL-no: 14.007] at a concentration of up to 0.4 \(\mu\text{mol/plate} (47 \mu\text{g/plate})\) (Sasagawa and Matsushima, 1991), isoquinoline [FL-no: 14.001] at a concentration of up to 50 \(\mu\text{g/mL}\), skatole [FL-no: 14.004] (3-methylindole) at a concentration of up to 0.4 \(\mu\text{mol/plate} (52 \mu\text{g/plate})\) (Szybalski, 1958; Sasagawa and Matsushima, 1991) or 3-acetylpyridine [FL-no: 14.039] at a concentration of up to 10,000 mg/plate (Pai et al., 1978).

In non-standardised assays, 2-acetylpyridine [FL-no: 14.038] at 0.50 to 0.87% (54,000 to 93,960 \(\mu\text{g/mL}\)) and 3-acetylpyridine [FL-no: 14.039] at 0.5 to 1.1% (55,100 to 122,322 \(\mu\text{g/mL}\)) caused a dose-dependent increase in mitotic aneuploidy in strain D61.M of *Saccharomyces cerevisiae* (Zimmermann et al., 1986). At the higher test concentrations, the growth of D61.M was strongly or completely inhibited. The authors noted that it is generally recognised that there is a threshold dose for induction of aneuploidy in yeast (Zimmermann et al., 1985a,b,c).

Assays in mammalian cell lines have been performed for isoquinoline [FL-no: 14.001] (Williams, 1984), skatole [FL-no: 14.004] (3-methylindole) (Kim et al., 1989) and pyrrole [FL-no: 14.041] (Williams, 1984). There was no evidence of increased unscheduled DNA synthesis when freshly isolated rat liver cells were incubated with pyrrole or isoquinoline (concentrations not specified) (Williams, 1984). Single-strand DNA breaks and inhibition of growth were reported when undeuterated or deuterated (at C2 or C3 positions) 3-methylindole (skatole) at 10 \(\mu\text{mol/L}\) to 1 mmol/L (1.31 to 131.18 \(\mu\text{g/mL}\)) was incubated with isolated cultured bovine kidney cells. However, there was no evidence of DNA interstrand crosslinks (Kim et al., 1989). These observations are consistent with reports that, at high concentrations, indoles deplete glutathione, leading to increased formation of DNA adducts (Nichols et al., 2000; Regal et al., 2001).

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\(^6\) The text is taken verbatim from the indicated reference source, but text related to substances not included in the present FGE has been removed.
In vivo

There was no evidence for mutation in a standard assay for sex-linked recessive lethal mutation when adult *Drosophila melanogaster* were fed 6-methylquinoline [FL-no: 14.042] at a concentration of 10 mmol/L (1,432 µg/mL) in a 5% sucrose solution for 3 days (Wild et al., 1983). Furthermore, 6-methylquinoline did not induce micronucleus formation in bone marrow cells obtained from male and female NMRI mice 30 h after treatment with the test compound as a single intraperitoneal dose at 0, 286, 429 or 572 mg/kg bw (Wild et al., 1983).

**JECFA's conclusion on genotoxicity**

Overall, negative results were reported in assays for reverse mutation in bacteria for six representative pyridine, pyrrole and quinoline derivatives (i.e. indole [FL-no: 14.007], isoquinoline [FL-no: 14.001], skatole [FL-no: 14.004] (3-methylindole), methyl 2-pyrrolyl ketone [FL-no: 14.047] (2-acetylpyrrole), pyrrole [FL-no: 14.041] and 3-ethylpyridine [FL-no: 14.061]). Although 6-methylquinoline gave positive results with metabolic activation, it gave negative results in studies *in vivo*, indicating that there are adequate detoxication mechanisms for the rapid absorption, distribution, biotransformation and elimination of the N-containing heteroaromatic derivatives. 2-Acetylpyrrole and 3-acetylpyridine produced positive results in yeast, but this is unlikely to occur at low doses because yeast is generally believed to have a threshold for the induction of aneuploidy. The positive results reported in bacteria for skatole (3-methylindole) are consistent with observations that, at high concentrations, indoles deplete glutathione, leading to reduced detoxification.

On the basis of the available evidence, the 22 pyridine, pyrrole and quinoline derivatives in this group do not demonstrate genotoxic potential.

For a summary of *in vitro/in vivo* genotoxicity data considered by JECFA, see Table A.1.

### 4.2.1.2. Genotoxicity studies – text taken from EFSA FGE.24Rev2 (EFSA CEF Panel, 2013)

**In vitro/in vivo**

Genotoxicity data were provided for seven of the 24 candidate substances. In *in vitro* studies on the candidate substances 2-methylindole [FL-no: 14.131], 2-methylpyridine [FL-no: 14.134], 3-methylpyridine [FL-no: 14.135], 4-methylpyridine [FL-no: 14.136], 2,4-dimethylpyridine [FL-no: 14.104], 3,5-dimethylpyridine [FL-no: 14.106] and 4-acetylpyridine [FL-no: 14.089] in doses up to 10000 µg/plate, with and without metabolic activation, did not cause reverse mutations in various strains of *S. Typhimurium* (Table A.2 in the present FGE.77Rev3).

Studies on induction of aneuploidy in *S. cerevisiae* D61.M available for the three candidate substances 2-methylpyridine [FL-no: 14.134], 2,4-dimethylpyridine [FL-no: 14.104] and 4-acetylpyridine [FL-no: 14.089] gave positive results. The positive results were obtained at high doses inhibiting the growth of the yeast. Furthermore, fungal systems for measuring aneuploidy have little relevance compared to the mammalian system.

No *in vivo* studies on genotoxicity of the candidate substances were available.

Genotoxicity tests are available for the eight supporting substances [FL-no: 14.004, 14.007, 14.038, 14.039, 14.041, 14.047, 14.061 and 14.065]. 2-Acetylpyrrole [FL-no: 14.047] (methyl 2-pyrrolyl ketone) was positive in TA98 without metabolic activation at the two highest concentrations tested. Negative results were obtained at the lowest concentration as well as with metabolic activation. This study is considered of limited relevance. Pyrrole [FL-no: 14.041], indole [FL-no: 14.007], 3-methylindole [FL-no: 14.004] (skatole), 3-ethylpyridine [FL-no: 14.061] and 2-acetylpyridine [FL-no: 14.038] were negative in bacterial mutation assays.

Studies on induction of aneuploidy in *S. cerevisiae* D61.M are available on three supporting substances, 2,6-dimethylpyridine [FL-no: 14.065], 2-acetylpyridine [FL-no: 14.038] and 3-acetylpyridine [FL-no: 14.039], which gave positive results. However, as for the three candidate substances, the positive results were obtained at high doses inhibiting the growth of the yeast. Furthermore, fungal systems for measuring aneuploidy have little relevance compared to the mammalian system.

**In vivo** data are available for one supporting substance.

3-Methylindole (skatole) [FL-no: 14.004] was reported negative in the MN assay in mice. The validity of this study, however, cannot be evaluated, as only an abstract is available.

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7 The text is taken verbatim from the indicated reference source, but text related to substances not included in the present FGE has been removed.
Positive results were obtained for some candidate and supporting substances in the Rec, DNA breaking, Chinese hamster ovary (CHO)\(^8\) and DNA synthesis assays. These results are, however, not considered valid.

**Conclusion on genotoxicity**

The genotoxicity data available for the candidate substances do not preclude their evaluation through the Procedure.

For a summary of in vitro/in vivo genotoxicity data considered by EFSA, see Tables A.2 and A.3 (Appendix A).

4.2.1.3. Genotoxicity studies evaluated by the panel in FGE.77Rev1

6-Methylquinoline [FL-no: 14.042]

6-Methylquinoline [FL-no: 14.042] was found to induce chromosome aberrations and sister chromatid exchanges (SCE) in CHO cells (National Toxicology Program (NTP), 1986).

A MN assay was performed by Nakajima (2005) essentially in line with the OECD Guideline 474. No significant increase of micronucleated polychromatic erythrocyte (PCE) frequency was observed in any groups of BDF1 male mice, treated by gavage at 225, 450 and 900 mg/kg bw for two subsequent days, 24 h apart. No significant decrease in the percentage of polychromatic erythrocytes to the analysed total erythrocytes (% PCE) was observed in any treatment group (Nakajima, 2005). The lack of cytotoxicity in the bone marrow cells does not allow a conclusion as to whether the test substance or a metabolite (e.g. an electrophilic epoxide) reached the bone marrow. Therefore, the results of this study have to be considered of limited relevance.

A bone marrow MN assay was performed by Honarvar (2004) on a structurally related substance, 6-isopropylquinoline, which was in compliance with good laboratory practice (GLP) and OECD test guideline 474. No significant increase of micronucleated PCE frequency was observed in any group of NMRI mice orally treated with 6-isopropylquinoline at 500, 1,000 and 2,000 mg/kg bw at 24 h after treatment and for the highest dose, 2,000 mg/kg bw also 48 h after treatment (Honarvar, 2004). Slight cytotoxic effects in the bone marrow (less than 10% changes in PCE/normochromatic erythrocyte (NCE) ratio) were observed, only at the high dose. Also, at the high dose group 48 h after treatment, the percentage of micronucleated cells (0.118) was higher than the corresponding vehicle control (0.065). The value was within the historical control range (up to 0.15%). Also in this case, due to the limited cytotoxicity, it is not clear whether the test substance/metabolite reached the target (bone marrow) in sufficient concentrations to elicit genotoxic effects.

The results of the MN assay by Honarvar (2004) on the structurally related substance 6-isopropylquinoline were considered of limited relevance due to the lack of evidence of target tissue exposure. Therefore, this study could not rule out the concern for genotoxicity for 6-methylquinoline.

For a summary of in vitro/in vivo genotoxicity data on 6-methylquinoline, see Tables A.4 and A.5 (Appendix A).

4.2.2. New genotoxicity studies evaluated in FGE.77Rev3

4.2.2.1. 6-Methylquinoline [FL-no: 14.042] – in vivo comet assay

6-Methylquinoline [FL-no: 14.042] (purity 99.9%) was tested for genotoxicity in vivo in male Han Wistar rats by assessing its ability to induce DNA damage in the liver and duodenum using the Comet assay (Keig-Shevlin, 2016). An additional separate experiment was conducted in the liver to clarify increases in tail intensity observed in the first experiment. The studies were in compliance with GLP and OECD test guideline 489 (OECD, 2014).

Doses were selected based on a range-finder experiment, where two doses of 6-methylquinoline [FL-no: 14.042] of 400 and 600 mg/kg bw per day were administered at 0 and 21 hrs to both male and female rats. No gender differences were observed for toxicity; therefore, only male rats were treated in the main experiments. The dose of 400 mg/kg bw per day was considered as an estimate of the maximum tolerated dose (MTD), based on clinical signs (ataxia, piloerection, decreased activity) observed. Based on this range-finder experiment, the following doses were tested in the main experiments: 0, 100, 200 and 400 mg/kg bw per day (six animals per group).

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\(^8\) This refers to a chromosomal aberration assay using CHO cells, where positive results were obtained at concentrations that show marked inhibition of DNA synthesis (Table A.2).
In the main experiments, 6-methylquinoline was administered twice to male Han Wistar rats by gavage at 0 (day 1) and 21 h (day 2). Vehicle (corn oil) and positive control (ethyl methanesulfonate (EMS), 200 mg/kg bw per day) groups were included. Tissues were sampled on day 2, 24 h after the first treatment.

No clinical signs of toxicity were observed following treatments with vehicle control, 6-methylquinoline or the positive control. There were no changes in clinical chemistry and no macroscopic or microscopic changes related to administration of 6-methylquinoline.

Experiment 1 – Comet assay in liver

Dose-related increases in group mean %tail intensity in the liver were observed in all test article treated groups. No dose-related increase in %clouds was observed. At the intermediate and high doses, the increases in tail intensity were statistically significant and a positive linear trend was also apparent (p ≤ 0.01). At 100 mg/kg bw per day, tail intensity and tail moment values were comparable with the vehicle control animals. At 200 mg/kg bw per day, a two-fold increase in group mean tail intensity and tail moment was observed. At 400 mg/kg bw per day, the group mean tail intensity and tail moment were three-fold higher than the vehicle control. The author of the study considered that this result was mainly due to two animals displaying elevated tail intensity and tail moment values which increased the group mean. As all individual animal data were within the laboratory’s historical control data and in the absence of any clinical chemistry or pathological changes in the liver which could be considered to be due to toxic effects of the test article, the biological relevance of these statistically significant increases in tail intensity was unclear. Therefore, the experiment in liver was repeated to check the reproducibility of these results.

The Panel considered the results of this experiment as positive taking into account that two criteria for evaluation and interpretation of results as positive (OECD test guideline 489) were fulfilled:

a) at least one of the test doses exhibits a statistically significant increase compared with the concurrent negative control,
b) the increase is dose related when evaluated with an appropriate trend test.

The third criterion ('any of the results are outside the distribution of the historical negative control data for a given species, vehicle, route, tissue, and number of administrations') is not applicable in this case because the range for historical negative controls is very wide (95% reference range of 0.08–5.08%).

Experiment 1 – Comet assay in duodenum

Following treatment with 6-methylquinoline at all dose levels, there was no evidence of any induction of DNA damage in cells isolated from duodenum. The group mean %tail intensity and tail moment values were comparable to the concurrent vehicle control group. No dose-related increase in %clouds in duodenum was reported following treatment with 6-methylquinoline.

Experiment 2 – Comet assay in liver

Group mean %tail intensity and tail moment values for all groups of animals treated with 6-methylquinoline were comparable with the group mean vehicle control data. There were no statistically significant differences in %tail intensity between treated and control groups. All individual animal data at all dose levels were generally consistent with the vehicle control animals and fell within the laboratory’s historical control data. There was no dose-related increase in %clouds in liver following treatment with 6-methylquinoline.

The increases in liver group mean tail intensity observed in experiment 1 were not reproduced in experiment 2. The authors of this study considered that the statistically significant increases in group mean liver tail intensity observed in the first experiment were due to chance occurrence in individual animals increasing the group means, rather than a true genotoxic effect and that there was no reproducible evidence that exposure of animals to 6-methylquinoline induced DNA damage in cells isolated from the liver.

Conclusion on comet assay

The Panel considered that the in vivo comet assay in rats (Keig-Shevlín, 2016) reported negative results in duodenum and equivocal results in liver, showing positive results in a first experiment, not reproduced in a second one. The Panel requested the applicant to clarify these results. The transgenic rodent gene mutation assay was considered as appropriate test for the in vivo follow-up of in vitro...
positive results in bacteria (EFSA Scientific Committee, 2011) In addition, a combined analysis of MN in peripheral blood in the same animals was requested to confirm the negative in vivo data observed in studies on chromosomal damage (Wild et al., 1983; Nakajima et al., 2005) which were considered by the Panel to be of only limited reliability.

4.2.2.2. 6-Methylquinoline [FL-no: 14.042] – in vivo gene mutation assay and micronucleus assay in Muta™ Mice

6-Methylquinoline (purity 99.9%) was tested for its ability to induce gene mutations in the lacZ transgene in liver and duodenum from treated male Muta™ Mice (CD2-lacZ80/HazfBR strain). In addition, an assessment of MN induction in peripheral blood reticulocytes from the same animals was included (Ballantyne, 2017).

Based on a range-finder experiment where no gender difference was observed, the MTD was established (500 mg/kg bw per day) for the main experiment based on clinical signs of toxicity (e.g. hunched posture, piloerection, semi-closed eyes, raised skin and fur, reduced activity and staggering). Groups of seven male mice were administered 6-methylquinoline, by oral gavage, at 0, 125, 250 or 500 mg/kg bw per day for 28 days. No positive control was included in the dosing regimen of this study, but was provided from another study carried out in the same laboratory measuring the same endpoints (positive control: ethylnitrosourea animals dosed at 10 mg/kg bw per day). The 28-day dosing period was followed by 3 days expression time before necropsy at day 31. On day 29, blood samples were collected from the tail vein of all the animals for the MN assay. After a 3-day expression period following the final dosing (day 31), the animals were sacrificed. All animals were macroscopically examined and the liver and duodenum were collected for the mutant frequency assay.

Mutant frequency in duodenum and liver

DNA was obtained from duodenum and liver tissues from all four treatment groups as well as frozen positive control DNA from animals treated with ethylnitrosourea from another study of the same laboratory for the mutant frequency assay. The DNA was packaged into phage heads ready to transfection in suspensions of E. coli, according to the OECD test guideline 488 (OECD, 2013). The transfected E. coli was plated, incubated and scored for plaques. According to the study report, all accepted packaging reactions resulted in at least 30,000 plaque-forming units (pfu) and at least one mutant plaque. All reported mutation data were generated for at least 200,000 pfu per tissue per animal, from at least three independent packaging reactions. At least 1 million pfu were obtained per group, per tissue from a minimum of five animals.

No statistically significant increases in mutant frequency were observed either in the liver or duodenum of treated Muta™ Mice, and all study results were within the laboratory’s historical control data set.

Micronucleus assay

Blood samples taken on day 29 were analysed for MN frequency by high speed flow cytometry. At least 20,000 reticulocytes from each sample were scored for MN.

No positive control was included in the MN experiment, but positive control blood samples (cyclophosphamide at 20 mg/kg bw per day) from another study of the same laboratory were analysed alongside the study samples. No significant increase in the frequency of micronucleated cells was observed in the peripheral blood reticulocytes of mice from any of the 6-methylquinoline treated groups compared to the vehicle controls. According to the author of the study report, all data fell within the historical control data set for the laboratory, but no historical control data were provided for animals dosed for more than seven days.

Conclusion on the in vivo gene mutation assay and in vivo micronucleus assay

The Panel concluded that under the conditions of this study, 6-methylquinoline did not induce mutations in the liver and duodenum of Muta™Mice, following treatment for 28 days up to the dose of 500 mg/kg bw per day (the MTD estimated for this study) and that the substance did not induce micronucleated cells in peripheral blood reticulocytes. In this study, no toxicity in liver and bone marrow was observed that would be considered as a direct evidence of tissue exposure to the tested substance. However, the clinical signs of toxicity observed in the dose range-finding study such as changes in behaviour (hunched posture, reduced activity and staggering), observed at high and mid doses) suggest that animals were systemically exposed to 6-methylquinoline, and this is considered as
a line of evidence of target tissue exposure (liver and bone marrow), on the basis of the criteria reported in the recent opinion of the EFSA Scientific Committee (EFSA Scientific Committee, 2017). The Panel considered this line of evidence, together with the observation that several other quinoline derivatives produce toxicity in liver and other tissue, sufficient to assume that 6-methylquinoline is systemically bioavailable and that, therefore, the negative results of the gene mutation assay in liver and the MN assay in peripheral blood are conclusive.

For a summary of in vivo genotoxicity data on 6-methylquinoline, see Appendix A, Table A.5.

4.2.2.3. EFSA considerations on genotoxicity

The Panel concluded that one of the 22 substances evaluated by JECFA, 6-methylquinoline [FL-no: 14.042], showed a genotoxic potential in vitro, with consistently positive results in several bacterial mutagenicity tests, after metabolic activation. Increases of chromosomal aberrations and sister chromatid exchanges in CHO cells were reported in a summary table (NTP, 1986), but were considered inadequate for the evaluation due to high level of toxicity at all the concentrations tested and the low number of metaphases examined.

6-Methylquinoline was reported negative in a test for gene mutations in Drosophila (Wild et al., 1983). Negative results were also reported in two in vivo MN studies in mice considered of limited relevance. In the first one (Wild et al., 1983), carried out by single ip administration, a single sampling time was used. In a more recent one, by gavage, no toxicity to the bone marrow was observed, although some weak evidence of systemic exposure of the animals was provided by clinical signs (reduction of spontaneous activity, prone position) reported at the highest dose tested (Nakajima, 2005).

The in vivo comet assay in rats reported negative results in duodenum and equivocal results in liver, showing a dose-related increase in %tail intensity in liver in a first experiment, which was not reproduced in a second one (Keig-Shevlin, 2016). A transgenic gene mutation assay in Muta™Mice (Ballantyne, 2017) carried out in liver and duodenum of animals treated with 6-methylquinoline for 28 days up to 500 mg/kg bw per day, the MTD, did not show any increase of mutant frequency. Therefore, the Panel considered that the concern for gene mutation is ruled out.

The analysis of MN in peripheral reticulocytes of the same animals after 28 days of treatment (Ballantyne, 2017) did not result in any increase in cells with MN compared to the control group. Therefore, the Panel considered that the concern for structural and numerical chromosomal aberrations is ruled out, likewise.

Based on the data available, the Panel concluded that there is no concern for genotoxicity of 6-methylquinoline [FL-no: 14.042]. For 21 JECFA evaluated pyridine, pyrrole and quinoline derivatives, the Panel concluded previously that the data available do not preclude evaluation through the Procedure. Based on the new data provided recently, the Panel concluded that also 6-methylquinoline [FL-no: 14.042] can now be evaluated through the Procedure.

4.2.3. Repeated dose toxicity studies

Since the publication of FGE.77 (EFSA, 2009), additional toxicity data have been provided for isoquinoline [FL-no: 14.001], pyrrole [FL-no: 14.041] and 2-acetylpyrrole [FL-no: 14.047], the latter also to cover the evaluation of the structurally related 2-propionylpyrrole [FL-no: 14.068]. The main studies provided are for each substance a 90-day study. Since the publication of FGE.77Rev1 (EFSA CEF Panel, 2014), additional toxicity data were provided for 1-furfurylpyrrole [FL-no: 13.134]. The main study provided is a 90-day study; these data are intended to cover the re-evaluation of this substance and 2-acetyl-1-ethylpyrrole [FL-no: 14.045] and 2-acetyl-1-methylpyrrole [FL-no: 14.046] and have been evaluated in FGE.77Rev2 (EFSA CEF Panel, 2015). In the present revision of FGE.77 (FGE.77Rev3), data from a 90-day study and from a carcinogenicity study on 6-methylquinoline [FL-no: 14.042] are evaluated.

The available toxicity studies are summarised in Appendix B, Table B.1.

4.2.3.1. Isoquinoline [FL-no: 14.001]

A 90-day oral study in rats was performed on isoquinoline according to the Japanese ‘Guidelines for designation of food additives and revision of standards for use of food additives, Notification No 29’ of the Environmental Health Bureau, Ministry of Health and Welfare, Japan, 22 March 1996. The
requirements of this guideline are very similar to the OECD Guideline 408. It is a GLP study. Groups (10/sex per dose) of male and female Sprague-Dawley rats were administered 0 (vehicle control), 0.03, 0.3 and 3 mg/kg bw per day of isoquinoline dissolved in corn oil by gavage daily for 90 and 91 days for males and females, respectively (Kojima, 2006). The purity of the test article was 98.5%. Animals were weighed at the start of the study and weekly thereafter. Food consumption and efficiency were measured weekly. The rats were caged individually during the experiment. All animals were subject to ophthalmologic examination prior to the start of the study, and on day 79, five animals of each sex per group were examined again. Urine was analysed on day 82 for five animals from each group. The rats were fasted for 18–21 h prior to blood sampling immediately prior to necropsy. A full haematological and biochemical analysis of blood was performed. At termination of the study, animals were sacrificed and subject to full necropsy. Histopathological examination was performed on all organs (as in the OECD Guideline 408) for the control and high dose group.

No animals died through the course of the study. No clinical signs of toxicity or behavioural changes were observed. Ophthalmological examination revealed no treatment-related changes. Mean body weights were comparable throughout the study between control and test groups of both sexes. Urine analysis did not reveal any treatment-related alterations when compared to controls. Haematology and blood chemistry results showed no significant differences between the test groups and controls. There were no organ weight changes or other macroscopic findings attributable to the administration of the test substance.

Histopathological examination did not show differences between controls and treated animals of either sex; some incidental findings occurred in both controls and treated animals, but there was no significant difference in their occurrence or intensity in the various organs when compared to the control groups.

Since there were no statistically significant changes due to the administration of the test material, the NOAEL of isoquinoline was determined to be 3 mg/kg bw per day in male and female rats after 90 days of administration by oral gavage (Kojima, 2006).

### 4.2.3.2. Pyrrole [FL-no: 14.041]

In a gavage study (Marumo, 2008), groups (10/dose per sex) of male and female Sprague-Dawley rats were administered 0 (vehicle control), 0.03, 0.3 and 3.00 mg/kg bw of aqueous pyrrole daily, by gavage for 90 days prior to necropsy. This study was performed according to ‘Guidelines for designation of food additives and for revision of standards for use of food additives’, Notification No. 29 of the Environmental Health Bureau, the Ministry of Health and Welfare, Japan, 22 March 1996 which is comparable to an OECD Guideline 408 study. It is a GLP study. Clinical observations were recorded daily and body weights and food consumption were recorded weekly. On day 79, five animals from each group were subject to ophthalmology examination. Urine samples were collected on day 82 for routine clinical chemical analysis. At termination, blood samples were taken for clinical chemistry determinations and haematological examination. At necropsy, organ weights for all organs required for an OECD Guideline 408 study were recorded. Tissues from all organs required in an OECD Guideline 408 study from both sexes of the control and 3.00 mg/kg bw per day groups were fixed and preserved for histopathological examination.

No mortality was observed throughout the course of the study and the general condition of the rats was unremarkable. Mean body weight gains and food consumption were comparable across test and control groups. Ophthalmologic examination revealed that in some animals in all male groups, controls included and some females of the 0.03 and 3.00 mg/kg bw per day groups, corneal clouding was observed. Urine analysis revealed no toxicologically significant findings except that one male rat out of five in the 3.00 mg/kg bw per day group showed some changes, suggesting a possible kidney effect at that level; however, there were no indications of kidney pathology in the histopathological findings of this rat. In the females, there were no effects observed in urinalysis except that they showed significantly higher concentrations of sodium, potassium and chloride ions, but this was not dose dependent. Males and females in the 3.00 mg/kg bw per day groups showed an increase in ‘urobilinogen concentrations’ in blood, but this was not accompanied by associated histopathology in the liver, spleen, bone marrow or haemolysis; the effect can be attributed to the interference of pyrrole present in urine in the colorimetric assay; it gives the same reaction as urobilinogen in the detection method used.

In female rats, the white blood cell count was lower for all three exposure levels than the control group, but this showed no dose relationship; the values were 4,600 ± 1,500, 3,600 ± 900, 3,400 ± 800 and 3,400 ± 1,100 per µL, respectively. At the two higher dose levels, this was
statistically significant (p < 0.05). None of the other haematological parameters was changed as compared to controls. In male rats, there was no difference in white blood cell levels or any other haematological parameter. Small changes in blood biochemical findings in male rats at the highest exposure were considered incidental.

Gross pathology examination revealed some organ weight variations including decreased absolute and relative pituitary gland weights only in low-dose treated male rats. All groups of male rats showed a somewhat decreased relative seminal vesicle weight due to a combination of increased body weight in the treated rats in combination with a slight decrease in absolute seminal vesicle weight. However, histopathology did not reveal abnormalities, neither in pituitary gland nor in seminal vesicles.

Histopathological examination was performed on all high dose and control animals, along with any tissues with lesions at other doses. In the lungs, alveolar accumulation of foamy cells was observed in eight males and three females at the 3.00 mg/kg bw dose and in four male controls. Mineralisation of the pulmonary arterial wall was reported for five males and two females of the high-dose group and two male controls. Focal thickening of alveolar septum with neutrophilic infiltration was seen in two high-dose male rats. Basophilic tubules were noted in the kidney cortex of eight males and five females of the high-dose group and five females of the control group. Atrophy of the seminiferous tubule was observed in two male in the high dose group, but the changes were very slight. In female in the high dose group, single animals showed follicle cysts or retention of the corpus luteum with a marked decrease of eosinophils in the endometrium and myometrium or marked mucification of the vaginal mucosa. Most of these phenomena were observed in both the treated and the control groups, and they are, therefore, considered incidental findings.

The lower white blood cell count in the females is considered an incidental finding and not considered an adverse effect since the count of all other blood cells types was normal in the female treated groups. In the males, no lower blood count for any cell types was observed, the histopathological examination revealed no correlating changes in the haematopoietic tissue and there was no dose–effect relationship (raising the question whether the control value was incidentally too high; the company, unfortunately, did not give an indication of historical control values in their report).

The Panel decided, based on the findings, that the NOAEL level was the highest exposure level 3.0 mg/kg bw per day.

4.2.3.3. 2-Acetylpyrrole [FL-no: 14.047]

A 14-day range finding study

In a 14-day range-finding dietary study (Bauter, 2012a), groups (3/sex per dietary intake level) of male and female Sprague-Dawley rats were fed a diet designed to provide 0 (dietary control), 1,000, 9,000 and 18,000 mg/kg feed of 2-acetylpyrrole [FL-no: 14.047] daily. These estimated dietary levels correspond to the measured intake of 0, 85, 550 and 842 mg/kg bw per day for males and 0, 91, 582 and 949 mg/kg bw per day for females, respectively. Individual food consumption was recorded on days 7 and 12. Due to increasing mortality in the high intake groups of both sexes, the study was terminated early at day 12. The results showed that the two higher doses were too toxic for a 90-day study. A 90-day study was started at lower exposure levels.

Effect on urinary iron excretion

The company also studied the effect of 2-acetylpyrrole [FL-no: 14.047] on urinary excretion of iron because 2-acetylpyrrole is a strong-complexing agent of metal ions. At a very high dose gavage study in rats (375 mg/kg bw orally for 10 days), the urinary excretion of total iron was increased six-fold (Mendes, 2012); no data are provided on absorption of iron from the intestinal tract, which might be influenced by complexation of iron with 2-acetylpyrrole.

90-day study

In an OECD test guideline 408 compliant 90-day study, groups of rats (10/sex per dietary intake level) of male and female Sprague-Dawley CD rats were fed a diet designed to provide 0 (dietary control), 1,050, 2,100 and 4,200 mg 2-acetylpyrrole [FL-no: 14.047]/kg feed daily (Bauter, 2012b). These dietary levels correspond to the calculated average daily intakes of 0, 68, 133 and 263 mg/kg bw for males and 0, 79, 155 and 298 mg/kg bw for females, respectively.

The test material was not stable in the diet, and in the report (Bauter, 2012b), it is suggested that part of it was probably not detected by the extraction method employed due to complexation with
metal ions in the feed. It is calculated that over the course of the study, the animals received concentrations of 35–40% of the target intake level on average. Therefore, values for exposure levels based on the measured intake are proportionally lower. Based on this analysis of the test diets, the mean daily intakes were calculated to be 367, 754 and 1,705 mg/kg feed. Assuming that the toxicity is only related to the free 2-acetylpyrrole, these dietary concentrations correspond to average daily intakes of 24, 48 and 107 mg/kg bw for males and 28, 56 and 121 mg/kg bw for females, respectively, over 90 days.

Clinical observations of toxicity were performed on day 0 and weekly until sacrifice. Animals were weighed on day 0 at the start of the study and weekly thereafter. Food consumption and efficiency were measured and calculated weekly. Blood chemistry and haematology were performed on blood drawn via sublingual bleed at day 43 for the controls and high intake groups and at day 86 for all groups after overnight fast. Urine was collected during the 15 h prior to the blood draw. Prior to initiation of the study and on day 91, the eyes of all rats were examined by focal illumination and indirect ophthalmoscopy. At termination of the study, all survivors were sacrificed and subject to full necropsy and histopathology as required by the OECD Guideline.

There were no mortalities or ophthalmological changes associated with the presence of 2-acetylpyrrole in the diet. Most other findings, generally also noted in control animals, were not considered adverse effects of test substance administration and were regarded as incidental. Statistically significant concentration-dependent reductions in body weight, body weight gain, food consumption (males and females) and food efficiency (females) at the highest dietary level (1,705 mg/kg feed measured concentration) during the study were attributed to the possible decrease in test substance palatability at high dietary levels.

Haematology parameters for both males and females were mostly unchanged during treatment. However, incidentally reaching a statistically significant difference when compared to concurrent controls, the values were in general within the range of historical controls and without associated histopathology correlate; they were, therefore, considered to be incidental and not related to the test material. However, statistically significantly (p < 0.05) decreased total white blood cell counts, erythrocyte counts, haemoglobin concentrations, haematocrit, absolute lymphocyte counts, absolute monocyte counts and absolute basophil counts and increased red cell distribution width were reported in the high intake group females on day 86. These parameters are outside of historical control levels although the variations are low in magnitude. There were no meaningful differences in coagulation parameters between test and control groups of both sexes.

Variations in clinical chemistry parameters were considered incidental and unrelated to the presence of 2-acetylpyrrole in the diet due to lack of concentration dependence or correlated pathology.

Organ weight measurements, absolute and relative brain weight, for males were comparable to controls, with some isolated exceptions; these were without histologic correlate and were considered unrelated to test substance in the diet.

Female rats of the high intake groups displayed minimal to moderate dark bilateral thyroid glands. Microscopic changes were slight thyroid hypertrophy/hyperplasia among 4/10 and 10/10 high intake group males and females, respectively. This was characterised by enlarged subgross tall columnar appearance of the follicular epithelial cells which appeared with fine cytoplasmic vacuolation with intermittent focally piled papillary projections into the follicular lumen. The company did not provide a clear (mechanistic) explanation for this finding.

In conclusion, although some haematology and clinical parameter changes were observed in mid- and high-dose groups, in the mid dose group were considered incidental and not of concern (not dose related and/or very small in magnitude and/or within historic controls and without histopathology correlation). However, the thyroid effects at the exposure level are of concern as well as the haematological changes in the high dose group females. Therefore, a NOAEL for 2-acetylpyrrole is derived from the middle dose 48 mg/kg bw per day in males and 56 mg/kg bw per day in females. The NOAEL value of 48 mg/kg bw per day is used in calculating the margin of safety.

Metabolites of 2-acetylpyrrole

Mendes (Mendes, 2012) analysed the urine obtained in metabolism cages from rats dosed with 2-acetylpyrrole [FL-no: 14.047] at 375 mg/kg by oral gavage as described above. Based on gas chromatography-mass spectroscopy (GC-MS) analysis, three major components were identified in the urine of both males and females treated with 2-acetylpyrrole. Unchanged 2-acetylpyrrole and pyrrol-2,5-dione were detected; the structure of another main metabolite detected in the urine is proposed to be 1,5-dihydropyrrol-2-one; however, further experiments have yet to be performed to confirm this.
4.2.3.4. 1-Furfurylpyrrole [FL-no: 13.134]

A 14-day range finding study

In a 14-day palatability and range-finding dietary study (Kappeler, 2013a), 1-furfurylpyrrole [FL-no: 13.134] was administered to male and female Crl:CD(SD) rats (3/sex per group) for 14 consecutive days in the diet to provide 0, 25, 75 and 200 mg/kg bw per day (Kappeler, 2013a). Observations for mortality and morbidity were performed twice daily and clinical examinations were performed once daily. Detailed physical examinations including body weight measurements were performed within 4 days of receipt, on the day of randomisation, prior to dosing on study day 0 (body weights, only) and weekly during the study. Food consumption was recorded 1 week prior to randomisation, on the day of randomisation and weekly throughout the study. Based on food consumption measurements, the calculated doses of 1-furfurylpyrrole were 0, 29, 84 and 211 mg/kg bw per day, respectively, for males and 0, 27, 81 and 192 mg/kg bw per day, respectively, for females over the entire study. All animals survived the 14 days of diet administration.

There were no overt adverse effects from 1-furfurylpyrrole administration. The only differences from control group animals were body weight reductions of 18% and 16% for males and females, respectively, in the 200 mg/kg bw per day group, that were associated with lower food consumption levels. It was concluded that 1-furfurylpyrrole was well tolerated in Crl:CD(SD) rats at dietary levels of up to 84 mg/kg bw per day and 81 mg/kg bw per day for males and females, respectively.

90-day study

In an OECD test guideline 408 compliant 90-day dietary study (GLP), 1-furfurylpyrrole [FL-no: 13.134] (purity 99.6%) was administered to individually housed Crl:CD(SD) rats (10/sex per group) at levels calculated to provide nominal doses of 0, 25, 75 and 150 mg/kg bw per day (Kappeler, 2013b). All animals were subject to observations for mortality and morbidity twice daily, daily clinical examinations and weekly detailed physical examinations, including body weight and food consumption measurements. Ophthalmic examinations were performed before the start of the study and in week 12. Clinical pathology parameters (haematology, coagulation, serum chemistry and urinalysis) were evaluated for all animals at the scheduled necropsy, after 13 weeks of treatment. Blood for haematology and serum chemistry was collected at necropsy after overnight fast and urine was collected overnight prior to necropsy. Coagulation was measured in blood collected from anaesthetised animals prior to sacrifice. All tissues were examined microscopically for animals of the 150 mg/kg bw per day groups. In addition, liver, spleen and nasal tissues were also examined for animals of the two lower intake level groups.

Based on food consumption data, the mean intake of 1-furfurylpyrrole was calculated to be 25, 77 and 154 mg/kg bw per day for males and 25, 75 and 151 mg/kg bw per day for females. All animals survived to the end of the study. There were no differences between treated and control groups noted in clinical observations, macroscopic findings or urinalysis, except for an increase in urobilinogen concentration.

By the time of necropsy, body weights were lower in the 150 mg/kg bw per day dose group for both males (−17%) and females (−11%) and in the 75 mg/kg bw per day dose group of females (−8%), relative to the control group. Decreased food consumption was also observed. These decreases were statistically significant for the male group at 150 mg/kg bw per day beginning on the first week and throughout the length of the study; for the female groups, the decrease was not statistically significant, although the weight gain in these females was decreased statistically significant. Slightly lower body weights were noted in the 25 and 75 mg/kg bw per day females and the 25 mg/kg bw per day females (4%, 5% and 6%, respectively).

Reductions in red blood cell counts (0.6%, 7.0* and 9.3**), haemoglobin levels (3.6%, 5.4* and 9.6**), and haematocrit values (3.7%, 7.2* and 9.9**) and slightly higher red blood cell distribution widths (4.1%, 4.9*** and 4.1***) were observed in all treated groups (25, 75 and 150 mg/kg bw per day, respectively) of males but were not observed in females. The effect at the lower exposure level was considered minor. Histopathology showed no changes in bone marrow at the highest dose level. Pigment deposits in the red pulp of the spleen were minimally increased only at the highest dose level in eight out of ten males; for females, this was one out of 10 at the middle dose and four out of 10 at the highest dose.

* significant at 0.05 according to Dunnett’s test.
** significant at 0.01 according to Dunnett’s test.
In males, dose-dependent changes in serum chemistry values included higher total bilirubin (0.01 ± 0.03, 0.02 ± 0.04, 0.07 ± 0.05** and 0.08 ± 0.04** mg/dL, respectively), which might be related to the reduced red blood cell counts and cholesterol (39*, 84** and 91***%, respectively). In females, similar increases were not dose dependent: 0.04 ± 0.05, 0.09 ± 0.03**, 0.11 ± 0.03*** and 0.10 ± 0.00** mg/dL for total bilirubin and 41**, 33* and 66***% for cholesterol. In parallel with the higher total bilirubin levels, the urobilinogen concentration in urine was increased at the higher dose levels only. At the highest dose, sorbitol dehydrogenase (SDH) values in both sexes were increased by 111** and 33% in males and females, respectively.

The increases in total bilirubin and cholesterol suggest effects on the liver. Higher mean liver weights relative to final body weight were noted in the male (6, 16** and 32**%) and female groups (12**, 17** and 32***% increase based on body weight). Centrilobular hepatocellular vacuolation was observed in the 75 and 150 mg/kg bw per day male groups and the 150 mg/kg bw per day group females only. The male groups scored 5/10 minimal and 5/10 mild at the highest dose; 4/10 minimal and 1/10 mild at the middle dose; 0/10 at the lowest dose; against 0/10 in the controls; the females showed only at the highest dose in 7/10 centrilobular vacuolisation against 0/10 in the control and both lower dose groups.

Other microscopic changes were identified in nasal sections and spleen. In the spleen, findings were limited to a very minimal increase in pigment deposits that were mainly restricted to the 150 mg/kg bw per day dose group. Microscopically, changes in nasal tissue included olfactory mucosal degeneration, depleted mucous secretion by goblet cells and thin deposits of a hyaline material along the surface of the olfactory epithelial cells. Olfactory mucosal degeneration was observed in both sexes at the 75 and 150 mg/kg bw per day intake levels and was slightly more prevalent and severe in males. The findings in the olfactory mucosa were considered to be adverse. Depleted mucous secretion by goblet cells was especially prominent in Nasal Section I and was observed in most rats administered 1-furfurylpyrrole.

It is concluded that exposure to 1-furfurylpyrrole [FL-no: 13.134] in the diet for 90 days resulted in adverse effects in the 75 and 150 mg/kg bw per day groups of males and females and consisted of effects on liver, red blood cells and olfactory mucosa. Therefore, the NOAEL of [FL-no: 13.134] was 25 mg/kg bw per day for male and female Crl:CD(SD) rats.

### 4.2.3.5. 6-Methylquinoline

6-Methylquinoline [FL-no: 14.042] was tested in a single-dose 90-day dietary study in Charles River CD rats (10–16 animals/sex per group) (Posternak et al., 1969). This publication is a summary of 90-day studies on 42 flavouring compounds. The publication states that ‘of these, 29 were at least 98% pure (..) the purity of the remainder was 90% or more’; the purity of 6-methylquinoline is not specified. The only available information on that study is a brief description of the method employed and a few general remarks. No study report or any details are available. The study results were submitted to a FEMA Expert Panel (before 1969).

A control and a single treatment group of the same sex were housed in pairs and given ad libitum access to water and food. The concentration of the test material in the diet was adjusted during the study to maintain a dietary intake of 2.2 mg/kg bw per day and 2.7 mg/kg bw per day in males and females, respectively. Food consumption and body weights were determined weekly. Limited haematology and blood urea analyses were conducted during the weeks 7 and 13 of the study, on half of the animals. At the end of the study, animals were sacrificed, liver and kidney weights were measured and ‘gross and histological examinations were carried out on a wide range of organs’.

The authors claim that at the exposure of 2.2 mg/kg bw per day in males and 2.7 mg/kg bw per day in females, 6-methylquinoline did not induce any effects in the parameters measured. However, the procedure of the 90-day study is very deficient compared to the current requirements for 90-day study according to OECD test guideline 408; moreover, the reporting is extremely deficient.

**Carcinogenicity study on 6-methylquinoline**

Fukushima et al. (1981) reported results from a 2-year oral carcinogenicity study on 6-methylquinoline in the rat. These data are part of a study which investigated the carcinogenicity of six quinoline derivatives including 8-nitroquinoline as positive control. In the publication by Fukushima et al. (1981), few study details are reported.

The compounds were mixed with the diet at 0.05%, except for the positive compound, 8-nitroquinoline, which was added at 0.10%. Thus, only one exposure level was tested in groups consisting of 43–46 animals per sex per group. Since pneumonia occurred during the last 12 weeks of
the experiment, many rats died prematurely. The surviving animals were necropsied after 104 weeks and ‘all organs’ were examined histologically. No details about the tissues examined are given.

In the 8-nitroquinoline-treated animals, extensive changes in the forestomach were observed in both males (n = 30) and females (n = 37): hyperplasia, papilloma and squamous cell carcinoma were seen in 97, 93 and 67%, respectively, of the males, and 97, 97 and 65% of the females. In addition, 30% (males) and 38% (females) had urinary bladder tumours and 8% of the females had a uterus carcinoma; in controls (n = 31 and 44 for males and females, respectively), the incidence of these tumours was 0%. Other tumours were of similar frequency as in controls.

For 6-methylquinoline (n = 38 and 37 for males and females, respectively, dose level: 500 mg/kg feed which is equivalent to 25 mg/kg bw per day), no differences with controls were observed; only the frequency of liver hyperplastic nodules seemed increased compared to controls: 10% vs. 0% in males and 13% vs. 7% in females.

These results show that the study did pick up signs of carcinogenicity for 8-nitroquinoline, while 6-methylquinoline was negative. The exposure level to these two substances was different: 0.10% in the feed for 8-nitroquinoline vs. 0.05% in the feed for 6-methylquinoline.

4.3. Other studies

Assay for tumour initiation

Quinoline and seven monomethylated derivatives, including 6-methylquinoline, were tested for tumour-initiating activity on SENCAR female mice skin with promotion by tetradecanoyl phorbol acetate (La Voie et al., 1984). The total initiation dose of either quinoline or the isomeric methylquinolines was 7.5 mg per mouse.

Quinoline induced tumours in 53% of the mice (on average 0.73 skin tumours per animal); a similar tumorigenic potential was observed for 4- and 8-methylquinoline (45% of the mice, with an average skin tumours per animal of 0.90 and 0.66, respectively). No statistically significant increases of skin tumours were observed for 2-, 3-, 5-, 6- and 7-methylquinoline.

These results suggest that 6-methylquinoline has no tumour-initiating properties in skin, under the conditions of this study. This would support the lack of carcinogenicity as reported in the study by Fukushima et al. (1981).

The Panel considered the data on 6-methylquinoline sufficient to disregard the observations with quinoline and other methylquinolines.

4.4. Application of the procedure

4.4.1. Application of the procedure to 22 pyridine, pyrrole and quinoline derivatives by JECFA (JECFA, 2005b, 2006a)

According to JECFA, three of the substances belong to structural class I, 13 to structural class II and six to structural class III, using the decision tree approach presented by Cramer et al. (1978).

JECFA concluded 20 pyridine, pyrrole and quinoline derivatives at step A3 in the JECFA Procedure, i.e. the substances are expected to be metabolised to innocuous products (step 2) and the intakes for all substances are below the thresholds for their structural classes I, II and III (step A3).

Two substances, 1-furfurylpyrrole [FL-no: 13.134] and 2-pyridine methanethiol [FL-no: 14.030], were evaluated via the B-side of the Procedure as the substances could not be anticipated to be metabolised to innocuous products. For these substances, the intake is below the threshold for the structural class III (step B3) and a NOAEL exists to provide an adequate margin of safety to the estimated intake as flavouring substances (step B4). For 1-furfurylpyrrole [FL-no: 13.134], a NOAEL of 12 mg/kg bw per day from a 90-day feeding study in rats (Morgareidge, 1971) is > 1,000,000 times greater than the estimated current intake of this substance as a flavouring substance. For 2-pyridine methanethiol [FL-no: 14.030], the NOAEL of 3.4 mg/kg bw per day from a 90-day feeding study in rats (Posternak et al., 1969) is > 20,000,000 times higher than the estimated current intake of this substance as a flavouring substance.

In conclusion, JECFA evaluated all 22 substances as to be of no safety concern at the estimated levels of intake as flavouring substances based on the MSDI approach.

The evaluations of the 22 pyridine, pyrrole and quinoline derivatives are summarised in Appendix C, Table C.1.
4.4.2. Application of the procedure to 24 pyridine, pyrrole, indole and quinoline derivatives from chemical group 28 evaluated by EFSA in FGE.24Rev2 (EFSA CEF Panel, 2013)

Twenty-four candidate substances were evaluated in FGE.24Rev2. Twenty-two of the 24 candidate substances are classified into structural class II and two substances into structural class III using the decision tree approach presented by Cramer et al. (1978).

Two of the substances, ethyl nicotinate [FL-no: 14.110] and isopropyl nicotinate [FL-no: 14.120], were concluded at step A3, i.e. the substances are expected to be metabolised to innocuous products (step 2) and the estimated daily intake is below the threshold for the structural class (step A3).

The remaining 22 substances were concluded at step B4, i.e. the substances could not be anticipated to be metabolised to innocuous products (step 2) and the estimated daily intake is below the threshold for the structural class (step B3). For the 22 substances, NOAELs could be derived to provide adequate margins of safety to the estimated levels of intake as favouring substance (step B4).

For the candidate substance 2-acetyl-5-methylpyrrole [FL-no: 14.085], a NOAEL of 48 mg/kg bw per day for the supporting substance 2-acetylpyrrole [FL-no: 14.047] is derived. The estimated daily per capita intake of 0.0012 \( \mu g \) for 2-acetyl-5-methylpyrrole [FL-no: 14.085] corresponds to 0.02 ng/kg bw per day at a body weight of 60 kg. Thus, a margin of safety of \( 2.4 \times 10^9 \) can be calculated. 2-Acetyl-5-methylpyrrole is accordingly not expected to be of safety concern at the estimated level of intake.

In an oral 37 weeks feeding study in rats on indole-3-carbinole, a substance structurally related to the two indole derivatives in this FGE (FGE.24Rev2), a NOAEL of 50 mg/kg bw per day could be derived. The combined estimated daily per capita intake of 0.0024 \( \mu g \) for 1-acetylindole [FL-no: 14.088] and 2-methylindole [FL-no: 14.131] corresponds to 0.04 ng/kg bw per day at a body weight of 60 kg. Thus, a margin of safety of \( 1.3 \times 10^9 \) can be calculated. 1-Acetylindole [FL-no: 14.088] and 2-methylindole [FL-no: 14.131] are accordingly not expected to be of safety concern at the estimated level of intake.

A 90-day oral feeding study in rats is available for the supporting substance 2-acetylpyridine [FL-no: 14.038]. The NOAEL derived is 37 mg/kg bw per day. The MSDI values for the 19 pyridine derivatives in this FGE (EFSA CEF Panel, 2013) are between 0.012 and 0.21 \( \mu g/capita \) per day. The combined estimated daily per capita intake of these 19 derivatives is 1.5 \( \mu g \), corresponding to 0.025 \( \mu g/kg \) bw per day. Thus, a margin of safety of \( 1.5 \times 10^6 \) can be calculated using the NOAEL of 37 mg/kg bw per day. The 19 pyridine derivatives in this flavouring group are accordingly not expected to be of safety concern at the estimated level of intake.

In conclusion, the Panel evaluated the 24 substances as to be of no safety concern at the estimated levels of intake as favouring substances based on the MSDI approach.

The stepwise evaluations of the 24 substances are summarised in Appendix C, Table C.2 of the present opinion.

4.4.3. EFSA consideration

The Panel agrees with the way the application of the Procedure has been applied by JECFA for four of the 22 substances. Methyl nicotinate [FL-no: 14.071], indole [FL-no: 14.007] and 3-methylindole [FL-no: 14.004] were evaluated via the A-side of the Procedure as they were anticipated to be metabolised to innocuous products. For these three substances, EFSA agreed no safety concern at step A3 of the Procedure, as the intake is below the threshold of the structural class (Cramer et al., 1978). 1-Furfurylpyrrole [FL-no: 13.134] and 2-pyridine methanethiol [FL-no: 14.030] were the only two substances evaluated through the B-side of the Procedure as the substances were not anticipated to be metabolised to innocuous products by JECFA. For 1-furfurylpyrrole [FL-no: 13.134], EFSA disagreed with JECFA, as the 90-day feeding study in rats (Morgareidge, 1971) was considered a poorly reported old study, the quality of which cannot be assessed, as stated in FGE.24 (EFSA, 2006). For 2-pyridine methanethiol [FL-no: 14.030], EFSA agrees with JECFA.

For 6-methylquinoline [FL-no: 14.042], contrary to JECFA, the Panel concluded in FGE.77, that this substance should not be evaluated using the Procedure until adequate \textit{in vivo} genotoxicity data become available. Additional genotoxicity data have become available for 6-methylquinoline after the publication of FGE.77, which have been evaluated in Revision 1 of FGE.77; however, the data were not sufficient to rule out the concern on the genotoxic potential of 6-methylquinoline. New \textit{in vivo} genotoxicity data have been provided for 6-methylquinoline, a Comet assay in duodenum and liver of

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9 [FL-no: 13.134] has been removed from FGE.24 Revision 2.
rats and a combined gene mutation assay and MN assay in MutaTM Mice. Based on these studies, the genotoxicity concern for 6-methylquinoline [FL-no: 14.042] can be ruled out.

The carcinogenicity study in rats on 6-methylquinoline (Fukushima et al., 1981) reports few study details and only one single dose of 6-methylquinoline was tested. Despite these limitations, this study suggests that 6-methylquinoline is not carcinogenic in rats under condition of the experiment. It induced an increase of liver hyperplastic nodules (not statistically significant) compared to controls. In addition, 6-methylquinoline did not induce skin tumours in a tumour-initiating test in mice (La Voie et al., 1984).

The only 90-day study in rats on 6-methylquinoline (Posternak et al., 1969) is of very limited quality, but was acceptable at the time the study was performed; it suggests that at a level around 2 mg/kg bw per day, no changes in a number of parameters measured occurred. If this dose is considered as a NOAEL, which is supported by the absence of strong indications of toxicity at higher exposure and longer duration in the study by Fukushima et al. (1981), an adequate margin of safety of approximately 400,000 can be calculated based on the MSDI.

For the remaining 16 substances, the Panel, in contrast to JECFA, did not anticipate that they will be metabolised to innocuous products due to concern with respect to N-oxidation of pyridines and for the pyrroles concerns about N-oxidation and epoxidation and accordingly concluded that they should be evaluated along the B-side. However, in FGE.77, for 10 [FL-no: 14.038, 14.039, 14.058, 14.059, 14.060, 14.061, 14.065, 14.066, 14.072 and 14.164] of these 16 substance, a NOAEL could be derived to provide adequate margins of safety to the estimated level of intakes as flavouring substance (step B4). A 90-day oral-feeding study in rats is available for 2-acetylpirydine [FL-no: 14.039]. The NOAEL derived is 37 mg/kg bw per day (Til and van der Meulen, 1971). The MSDI values for the 10 pyridine derivatives in this FGE are between 0.06 and 50 μg/capita per day. The combined estimated daily per capita intake of the 10 pyridine derivatives evaluated through the B-side is 57 μg corresponding to 0.95 μg/kg bw per day. Thus, a margin of safety of approximately 39,000 can be calculated using the NOAEL of 37 mg/kg bw per day. The 10 pyridine derivatives in this flavouring group evaluated through the B-side are accordingly not expected to be of safety concern at the estimated levels of intake.

For pyrrole [FL-no: 14.041] and the five pyrrole derivatives [FL-no: 13.134, 14.045, 14.046, 14.047 and 14.068] as well as for isoquinoline [FL-no: 14.001], NOAELs could not be derived as such or for structurally related substances in FGE.77. Accordingly, additional toxicological data were required for these seven substances (step B4) in FGE.77.

Additional toxicity data have become available, after the publication of FGE.77, for isoquinoline [FL-no: 14.001], pyrrole [FL-no: 14.041] and 2-acetylpyrrole [FL-no: 14.047] (Kojima, 2006, Marumo, 2008 and Bauter, 2012b), the latter also to cover the evaluation of the structurally related 2-propionylpyrrole [FL-no: 14.068].

Based on the data for isoquinoline [FL-no: 14.001] (Kojima, 2006), a NOAEL of 3 mg/kg bw per day could be established. When comparing this NOAEL at step B4 in the Procedure to the estimated exposure based on the MSDI (0.012 μg/capita per day, corresponding to 0.0002 μg/kg bw per day), an adequate margin of safety of 15 × 10⁵ can be calculated.

Based on the data for pyrrole [FL-no: 14.041] (Marumo, 2008), a NOAEL of 3 mg/kg bw per day could be established. When comparing this NOAEL at step B4 in the Procedure to the estimated exposure based on the MSDI (0.11 μg/capita per day, corresponding to 0.0018 μg/kg bw per day), an adequate margin of safety of 16 × 10⁵ can be calculated.

Based on the data for 2-acetylpyrrole [FL-no: 14.047] (Bauter, 2012b), a NOAEL of 48 mg/kg bw per day could be established. When comparing the NOAEL at step B4 in the Procedure to the estimated exposure based on the MSDI (3.3 μg/capita per day, corresponding to 0.055 μg/kg bw per day), an adequate margin of safety of 87 × 10⁵ can be calculated. For 2-propionylpyrrole [FL-no: 14.068], supported by 2-acetylpyrrole [FL-no: 14.047], the MSDI is 0.012 μg/capita per day, which is well below the MSDI of 2-acetylpyrrole and accordingly not expected to be of safety concern at the estimated levels of intake.

Additional toxicity data have become available, after the publication of FGE.77Rev1, for 1-furfurylpyrrole [FL-no: 13.134] (Kappeler, 2013b). The data are intended to cover the re-evaluation of this substance and 2-acetyl-1-ethylpyrrole [FL-no: 14.045] and 2-acetyl-1-methylpyrrole [FL-no: 14.046]. The main study provided is a 90-day study.

Based on the data submitted for 1-furfurylpyrrole [FL-no: 13.134] (Kappeler, 2013b), a NOAEL of 25 mg/kg bw per day could be established. When comparing this NOAEL at step B4 in the Procedure to the estimated exposure based on the MSDI (0.12 μg/capita per day, corresponding to 0.002 μg/kg bw per day), an adequate margin of safety of 12.5 × 10⁵ can be calculated. For 2-acetyl-1-ethylpyrrole [FL-no: 14.045], supported by 1-furfurylpyrrole [FL-no: 13.134], the MSDI is also 0.12 μg/
capita per day and accordingly not expected to be of safety concern at the estimated levels of intake. For 2-acetyl-1-methylpyrrole [FL-no: 14.046], the MSDI is 1.2 μg/capita per day which is 10 times the figure for the MSDI of 1-furfurylpyrrole resulting in an adequate margin of safety of $12.5 \times 10^5$; 2-acetyl-1-methylpyrrole [FL-no: 14.046] is accordingly not expected to be of safety concern at the estimated levels of intake.

5. Conclusions

The present Revision of FGE.77, FGE.77Rev3, includes the assessment of additional genotoxicity data for 6-methylquinoline [FL-no: 14.042]. The studies provided are an in vivo Comet assay in rats and a combined gene mutation and MN assay in Muta\textsuperscript{TM}Mice. The applicant also submitted toxicity data for 6-methylquinoline.

The Panel concluded that the 22 substances in the JECFA flavouring group of pyridine, pyrrole and quinoline derivatives are structurally related to the group of pyridine, pyrrole, indole and quinoline derivatives from chemical group 28 evaluated by EFSA in the Flavouring Group Evaluation 24, Revision 2 (FGE.24Rev2).

JECFA evaluated two substances [FL-no: 13.134 and 14.030] via the B-side of the Procedure and 20 substances via the A-side.

The Panel agrees with the way that the application of the Procedure has been applied by JECFA for four of the 22 substances. Three of these four substances, methyl nicotinate [FL-no: 14.071], indole [FL-no: 14.007] and 3-methylindole [FL-no: 14.004], were evaluated by JECFA on the A-side of the Procedure, as they were anticipated to be metabolised to innocuous products. For these three substances, the Panel agreed no safety concern at step A3 of the Procedure, as their intake estimates (MSDI) were below the threshold of their structural class. For the fourth substance, 2-pyridine methanethiol [FL-no: 14.030], the Panel agreed with JECFA that it should be evaluated through the B-side of the Procedure, as the substance was not anticipated to be metabolised to innocuous products. However, a NOAEL was derived from a 90-day study, which provided an adequate margin of safety.

For 1-furfurylpyrrole [FL-no: 13.134], EFSA disagreed with JECFA, as the 90-day feeding study in rats was considered a poorly reported old study, the quality of which cannot be assessed. However, for this substance, additional toxicity data were submitted and evaluated in FGE.77Rev2. Based on these data, adequate margins of safety could be calculated for [FL-no: 13.134, 14.045 and 14.046].

For 16 substances, the Panel, in contrast to JECFA, did not anticipate that they will be metabolised to innocuous products and accordingly concluded that they should be evaluated along the B-side of the Procedure. However, in FGE.77, for 10 [FL-no: 14.038, 14.039, 14.058, 14.059, 14.060, 14.061, 14.065, 14.066, 14.072 and 14.164] of these 16 JECFA-evaluated pyridine derivatives evaluated via the B-side of the Procedure by EFSA, NOAELs could be derived to provide adequate margins of safety and the Panel agrees with the JECFA conclusion 'no safety concern at estimated levels of intake as flavouring substances' based on the MSDI approach.

In FGE.77Rev1, three 90-day studies were evaluated for isoquinoline [FL-no: 14.001], pyrrole [FL-no: 14.041] and 2-acetylpyrrole [FL-no: 14.047] and NOAELs to provide adequate margin of safety were derived to cover these three substances as well as the structurally related 2-propionylpyrrole [FL-no: 14.068].

In FGE.77Rev2, one 90-day study was evaluated for 1-furfurylpyrrole [FL-no: 13.134]. A NOAEL to provide adequate margin of safety was derived to cover this substance as well as the structurally related 2-acetyl-1-ethylpyrrole [FL-no: 14.045] and 2-acetyl-1-methylpyrrole [FL-no: 14.046].

In previous revisions of this FGE, the Panel concluded, contrary to JECFA, that 6-methylquinoline [FL-no: 14.042] (evaluated via the A-side by JECFA) should not be evaluated through the Procedure due to concern with respect to genotoxicity in vitro.

Based on the available data and the new in vivo genotoxicity studies on 6-methylquinoline [FL-no: 14.042], the Panel concluded that, for 6-methylquinoline [FL-no: 14.042], there is no concern with respect to genotoxicity. Therefore, 6-methylquinoline [FL-no: 14.042] can now be evaluated through the Procedure.

The carcinogenicity study in rats on 6-methylquinoline (Fukushima et al., 1981) reports few study details and only one single dose (0.05% in diet which is equivalent to 25 mg/kg bw per day) of 6-methylquinoline was tested. Despite these limitations, this study suggests that 6-methylquinoline is not carcinogenic in rats under the condition of the experiment. In addition, 6-methylquinoline did not induce skin tumours in a tumour-initiating test in mice (La Voie et al., 1984).

The only 90-day study in rats on 6-methylquinoline (Posternak et al., 1969) is of very limited quality, but was acceptable at the time the study was performed; it suggests that at a level around 2 mg/kg bw
per day, no changes in a number of parameters measured occurred. If this dose is considered as a NOAEL, an adequate margin of safety of approximately 400,000 can be calculated based on the MSDI.

From the group of 22 substances evaluated by JECFA, the Panel evaluated three substances via the A-side [FL-no: 14.004, 14.007, 14.071] and 19 substances via the B side: [FL-no: 13.134, 14.001, 14.030, 14.038, 14.039, 14.041, 14.042, 14.045, 14.046, 14.047, 14.058, 14.059, 14.060, 14.061, 14.065, 14.066, 14.068, 14.072 and 14.164]. For these 19 substances, NOAELs could be derived, which provided adequate margins of safety.

For the following substances [FL-no: 13.134, 14.001, 14.030, 14.041, 14.042, 14.058, 14.072], the Industry has submitted use levels for normal and maximum use. For the remaining 15 substances, use levels are needed to calculate the mTAMDIs in order to identify those flavouring substances that need more refined exposure assessment and to finalise the evaluation.

In order to determine whether the conclusion for the 22 JECFA evaluated substances can be applied to the materials of commerce, it is necessary to consider the available specifications. Adequate specifications including complete purity criteria and identity tests are available for the 22 JECFA-evaluated substances.

Thus, for the 22 JECFA evaluated pyridine, pyrrole and quinoline derivatives [FL-no: 13.134, 14.001, 14.004, 14.007, 14.030, 14.038, 14.039, 14.041, 14.042, 14.045, 14.046, 14.047, 14.058, 14.059, 14.060, 14.061, 14.065, 14.066, 14.068, 14.071, 14.072 and 14.164], the Panel agrees with the JECFA conclusion ‘no safety concern at estimated levels of intake as flavouring substances’ based on the MSDI approach.

**Documentation provided to EFSA**

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Abbreviations

bw Body Weight
CAS Chemical Abstract Service
CEF Panel on Food Contact Materials, Enzymes, Flavourings and Processing Aids
CHO Chinese hamster ovary (cells)
CoE Council of Europe
DNA Deoxyribonucleic acid
EPA United States Environmental Protection Agency
FAO Food and Agriculture Organization of the United Nations
FEMA Flavor and Extract Manufacturers Association
FGC Flavouring Group Evaluation
FLAVIS (FL) Flavour Information System (database)
GLP Good Laboratory Practise
GC-MS Gas chromatography-mass spectrometry
ID Identity
i.p. Intraperitoneal
IR Infrared spectroscopy
JEFCFA The Joint FAO/WHO Expert Committee on Food Additives
MN micronucleus
MSDI Maximised Survey-derived Daily Intake
mTAMDI Modified Theoretical Added Maximum Daily Intake
NCE Normochromatic erythrocyte
No Number
NOAEL No observed adverse effect level
NTP National Toxicology Program
OECD Organisation for Economic Co-operation and Development
PCE Polychromatic erythrocyte
PCR Polymerase Chain Reaction
pfu plaque-forming units
SCE Sister chromatic exchange
SCF Scientific Committee on Food
SDH sorbitol dehydrogenase
WHO World Health Organization
## Appendix A – Summary of genotoxicity data

### Table A.1: Genotoxicity data (in vitro/in vivo) JECFA (JECFA, 2006a)

| FL-no | EU Register name | Structural formula | End-point | Test system | Concentration | Results | Reference |
|-------|------------------|--------------------|-----------|-------------|---------------|---------|-----------|
| 14.007 1301 | Indole | ![Indole structural formula](image) | Reverse mutation | S. Typhimurium TA100 | ≤ 20 µg/plate | Negative<sup>(a)</sup> | Ochiai et al. (1986) |
| | | | Reverse mutation | S. Typhimurium TM677 | 4 mmol/L (469 µg/mL)<sup>(b)</sup> | Negative<sup>(c)</sup> | Kaden et al. (1979) |
| | | | Reverse mutation | S. Typhimurium TA98, TA100, TA1535, TA1538 | 4-2,500 µg/plate | Negative<sup>(d)</sup> | Anderson and Styles (1978) |
| | | | Reverse mutation | S. Typhimurium TA98, TA100 | ≤ 500 nmol/plate (59 µg/plate)<sup>(b)</sup> | Negative<sup>(a)</sup> | Vance et al. (1986) |
| | | | Reverse mutation | S. Typhimurium TA100, TA1535, TA1537 | 3 µmol/plate (351 µg/plate)<sup>(b)</sup> | Negative<sup>(d)</sup> | Florin et al. (1980) |
| | | | Reverse mutation | S. Typhimurium TA98 | 0.03-30 µmol/plate (3.5-3515 µg/plate)<sup>(b,e)</sup> | Negative<sup>(d)</sup> | Florin et al. (1980) |
| | | | Reverse mutation | S. Typhimurium TA97, TA102 | 10–1,000 µg/plate | Negative<sup>(d)</sup> | Fujita et al. (1994) |
| | | | Reverse mutation | S. Typhimurium TA98, TA100 | ≤ 0.4 µmol/plate (47 µg/plate)<sup>(b)</sup> | Negative<sup>(d)</sup> | Sasagawa and Matsushima (1991) |
| | | | Mutation | E. coli WP2 uvrA/pKM101 | ≤ 0.4 µmol/plate (47 µg/plate)<sup>(b)</sup> | Negative<sup>(d)</sup> | Sasagawa and Matsushima (1991) |
| FL-no | EU Register name | Structural formula | End-point | Test system | Concentration | Results | Reference |
|-------|-----------------|-------------------|-----------|-------------|---------------|---------|-----------|
| FL-no | JECFA-no | JECFA name | | | | |
| 14.042 | 1302 | 6-Methylquinoline | | | | |
| Reverse mutation | S. Typhimurium TA100 | 100–600 μg/plate | Positive(c) | Dong et al. (1978) |
| Reverse mutation | S. Typhimurium TA98, TA100 TA1535, TA1537 and TA1538 | $\leq$ 3,600 μg/plate | Negative(a), Positive(c,f) | Wild et al. (1983) |
| Reverse mutation | S. Typhimurium TA98 and TA100 | $\leq$ 6 μmol/plate (859 μg/plate)(g) | Negative(a) Positive(c) | Nagao et al. (1977) |
| Reverse mutation | S. Typhimurium TA98 and TA100 | $\leq$ 1,000 μg/plate | Negative(a) Positive(c) | Zeiger et al. (1992) |
| Reverse mutation | S. Typhimurium TA98 and TA100 | NR | Negative(a) Positive(c) | Sugimura et al. (1976) |
| Reverse mutation | S. Typhimurium TA100 | 5 μmol/plate (716 μg/plate)(g) | Positive(c) | Takahashi et al. (1988) |
| Reverse mutation | S. Typhimurium TA98 | NR | Negative(d) | Debnath et al. (1992) |
| Reverse mutation | S. Typhimurium TA100 | 3.3–333 μg/plate | Negative(a) Positive(c) | Debnath et al. (1992) |
| 14.001 | 1303 | Isoquinoline | | | | |
| Reverse mutation | S. Typhimurium TA98 and TA100 | 20–50 μg/mL | Negative(d) | Sideropoulos and Specht (1984) |
| Reverse mutation | S. Typhimurium TM677 | $\leq$ 8 mmol/L (1,033 μg/mL)(h) | Negative(c) | Kaden et al. (1979) |
| Reverse mutation | S. Typhimurium TA98 and TA100 | NR | Negative(d) | Sugimura et al. (1976) |
| Reverse mutation | S. Typhimurium TA98 and TA100 | 1–20 μmol/plate (129–2,583 μg/plate)(h) | Negative(d) | Nagao et al. (1977) |
| Reverse mutation | S. Typhimurium TA98 and TA100 | 10,000–20,000 μg/mL | Negative(d) | Epler et al. (1979) |
| Mutation | E. coli B/r HCR+ | 50 μg/mL | Negative(d) | Sideropoulos and Specht (1984) |
| Unscheduled DNA synthesis | Rat hepatocytes | NR | Negative | Williams (1984) |
| FL-no | EU Register name | Structural name | Structural formula | End-point | Test system | Concentration | Results | Reference |
|-------|------------------|------------------|-------------------|-----------|-------------|---------------|---------|-----------|
| 14.004| 3-Methylindole   |                  | ![3-Methylindole](image) | Reverse mutation | S. Typhimurium TA100, TA1535 and TA1537 | 3 μmol/plate (394 μg/plate) | Negative | Florin et al. (1980) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA98 | 0.03–30 μmol/plate (3.9–3,935 μg/plate) | Negative | Florin et al. (1980) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA98 and TA100 | NR | Negative | Kim et al. (1989) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA98 and TA100 | ≤ 0.4 μmol/plate (52 μg/plate) | Negative | Sasagawa and Matsushima (1991) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA100 | ≤ 100 μg/plate | Negative | Ochiai et al. (1986) |
|  |                  |                  |                   | Mutation | E. coli WP2 uvrA/pKM101 | ≤ 0.4 μmol/plate (52 μg/plate) | Negative | Sasagawa and Matsushima (1991) |
|  |                  |                  |                   | Mutation | E. coli Sd-4-73 | 0.01–0.025 mL/disk | Negative | Szybalski (1958) |
|  |                  |                  |                   | DNA single-strand break | Bovine kidney cells | 10 μmol/1 mmol/L (1.31–131.18 μg/mL) | Positive | Kim et al. (1989) |
| 14.047| 2-Acetylpyrrole  |                  | ![2-Acetylpyrrole](image) | Reverse mutation | S. Typhimurium TA98 and TA100 | 12.5–200 μg/plate | Negative | Wang et al. (1994) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA98 | 4–100 μmol/plate (437–10,913 μg/plate) | Negative | Lee et al. (1994) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA100 | 4–100 μmol/plate (437–10,913 μg/plate) | Positive | Lee et al. (1994) |
| 14.038| 2-Acetylpyridine |                  | ![2-Acetylpyridine](image) | Mitotic aneuploidy | S. cerevisiae D61.M | 0.50–0.87% (54,000–939,600 μg/mL) | Positive | Zimmermann et al. (1986) |
| 14.041| Pyrrole          |                  | ![Pyrrole](image)  | Reverse mutation | S. Typhimurium TA98, TA100 and TA102 | 14 nmol/plate (0.94–9,926 μg/plate) | Negative | Aeschbacher et al. (1989) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA100, TA1535 and TA1537 | 3 μmol/plate (201 μg/plate) | Negative | Florin et al. (1980) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA98 | 0.03–30 μmol/plate (2.01–2013 μg/plate) | Negative | Florin et al. (1980) |
|  |                  |                  |                   | Reverse mutation | S. Typhimurium TA98 and TA100 | NR | Negative | Lee et al. (1994) |
|  |                  |                  |                   | Unscheduled DNA synthesis | Rat hepatocytes | NR | Negative | Williams (1984) |
| FL-no | EU Register name | Structural formula | End-point | Test system | Concentration | Results | Reference |
|-------|------------------|--------------------|-----------|-------------|---------------|---------|-----------|
| 14.061 1315 | 3-Ethylpyridine | | Reverse mutation | S. Typhimurium TA98, TA100, TA1535 and TA1537 | 3 μmol/plate (321 μg/plate) | Negative<sup>(d)</sup> | Florin et al. (1980) |
| 14.039 1316 | 3-Acetylpyridine | | Mutation | E. coli WP2 uvrA | 5,000–10,000 μg/plate | Negative | Pai et al. (1978) |
| | | | Mitotic aneuploidy | S. cerevisiae D61.M | 0.5–1.11% (55,100–122,322 μg/mL) | Positive | Zimmermann et al. (1986) |

**In vivo**

| In vivo FL-no | EU Register name | Structural formula | End-point | Test system | Concentration | Results | Reference |
|--------------|------------------|--------------------|-----------|-------------|---------------|---------|-----------|
| 14.042 1302 | 6-Methylquinoline | | Sex-linked recessive mutation | Drosophila melanogaster | 10 mmol/L (1432 μg/mL)<sup>(g)</sup> | Negative | Wild et al. (1983) |
| | | | Micronucleus formation | NMRI mouse | 0, 286, 429 or 572 mg/kg bw | Negative | Wild et al. (1983) |

NR: Not reported.

(a): Without metabolic activation.
(b): Calculated based on relative molecular mass = 117.15.
(c): With metabolic activation.
(d): With and without metabolic activation.
(e): Toxic at concentrations > 3.0 mmol/plate (351 mg/plate).
(f): TA100 and TA1535.
(g): Calculated based on relative molecular mass = 143.19.
(h): Calculated based on relative molecular mass = 129.16.
(i): Calculated based on relative molecular mass = 131.18.
(j): Toxic at concentrations of > 3.0 mmol/plate (394 mg/plate).
(k): Calculated based on relative molecular mass = 109.13.
(l): Calculated based on density = 1.08 g/mL (Sigma-Aldrich, 2003; available at http://www.sigmaaldrich.com).
(m): Calculated based on relative molecular mass = 67.09.
(n): Calculated based on density = 1.102 g/mL (Sigma-Aldrich, 2003; available at http://www.sigmaaldrich.com).
### Table A.2: Genotoxicity data (in vitro) EFSA/FGE.24Rev2 (EFSA CEF Panel, 2013)

| Chemical Name [FL-no] | Test System | Test Object | Concentration | Result | Reference | Comments |
|-----------------------|-------------|-------------|---------------|--------|-----------|---------|
| (Pyrrole [14.041])    | Ames assay  | *S. Typhimurium* TA98; TA100; TA102 | 1.4 mmol/plate (93,926 µg/plate) | Negative<sup>(a)</sup> | Aeschbacher et al. (1989) |         |
|                       | Ames assay  | *S. Typhimurium* TA100; TA1535; TA1537 | 3 µmol/plate (201 µg/plate) | Negative<sup>(a)</sup> | Florin et al. (1980) |         |
|                       | Ames assay  | *S. Typhimurium* TA98 | 30 µmol/plate (2013 µg/plate) | Negative<sup>(a)</sup> | |         |
|                       | Ames assay  | *S. Typhimurium* TA98; TA100 | | Not reported | Lee et al. (1994) |         |
| Rec assay             | *B. subtilis* H17 (rec<sup>+</sup>), M45 (rec<sup>−</sup>) | 4 and 20 mg/disk | Positive<sup>(c)</sup> | Kim et al. (1987) | Poor predictive value for mutagenicity. Limited value |
| Unscheduled DNA synthesis | Rat hepatocytes | | Negative | Williams (1984) |         |
| 1-Methylpyrrole       | Ames assay  | *S. Typhimurium* TA98; TA100; TA102 | 11 nmol-1.1 mmol/plate | Negative<sup>(a)</sup> | Aeschbacher et al. (1989) | 6 dose levels. The study is considered valid |
|                       | Rec assay   | *B. subtilis* H17 (rec<sup>+</sup>) M45 (rec<sup>−</sup>) | 2, 4, 20 and 40 mg/disk (500.5 µmol/disk) | Positive<sup>(a)</sup> | Kim et al. (1987) | Poor predictive value for mutagenicity. Limited value |
| Chemical Name [Fl-no] | Test System | Test Object | Concentration | Result | Reference | Comments |
|-----------------------|-------------|-------------|---------------|--------|-----------|----------|
| (Indole [14.007])     | Ames assay  | *S. Typhimurium* TA100 | 20 μg/plate | Negative(b) | Ochiai et al. (1986) |         |
| Ames assay            | *S. Typhimurium* TM677 | 4 mM (469 μg/mL) | Negative(c) | Kaden et al. (1979) |         |
| Ames assay (plate incorporation method) | *S. Typhimurium* TA98; TA100; TA1535; TA1538 | 2,500 μg/plate | Negative(a) | Anderson and Styles, (1978) |         |
| Ames assay            | *S. Typhimurium* TA98; TA100 | 500 nmol/plate (59 μg/plate) | Negative(b) | Vance et al. (1986) |         |
| Ames assay (preincubation method) | *S. Typhimurium* TA100; TA1535; TA1537 | 3 μmol/plate (351 μg/plate) | Negative(a) | Florin et al. (1980) |         |
|                       | *S. Typhimurium* TA98 | 30 μmol/plate (3,515 μg/plate) | Negative(c) |         |           |
|                       | *S. Typhimurium* TA97; TA102 | 1,000 μg/plate | Negative(a) | Fujita et al. (1994) |         |
|                       | *S. Typhimurium* TA98; TA100 *E. coli* WP2uvrA/pKM101 | 0.4 μmol/plate (47 μg/plate) | Negative(a) | Sasagawa and Matsushima (1991) |         |
|                       | *S. Typhimurium* TA100 | 500 μg/plate | Negative(b) | Hashizume et al. (1991) |         |
| 2-Methylindole [14.131] | Ames assay (preincubation method) | *S. Typhimurium* TA98; TA100; TA1535; TA1538 | 4, 20, 100, 500 and 2,500 μg/plate | Negative(a) | Anderson and Styles (1978) | The study is considered valid |
|                       | *S. Typhimurium* TA98; TA100; TA1535; TA1537 | 3 μmol/plate (394 μg/plate) | Negative(a) | Florin et al. (1980) | Single-dose study |
| Ames assay            | *S. Typhimurium* TA98 | 3 nmol·30 μmol/plate (12 doses) (3,935 μg/plate) | Negative(a) | Curvall et al. (1982) | The study is considered valid. |
|                       | *S. Typhimurium* TM677 | 2 mM (262 μg/mL) | Negative | Kaden et al. (1979) | Single-dose study. |
| Chemical Name                  | Test System                          | Test Object         | Concentration                  | Result     | Reference                       | Comments                                                                                   |
|-------------------------------|--------------------------------------|---------------------|--------------------------------|------------|---------------------------------|-------------------------------------------------------------------------------------------|
| (3-Methylindole [14.004])     | Ames assay (preincubation method)    | S. Typhimurium TA100; TA1535; TA1537 | 3 μmol/plate (394 μg/plate)  | Negative(a) | Florin et al. (1980)             |                                                                                           |
|                               |                                      | S. Typhimurium TA98  | 30 μmol/plate (3,935 μg/plate) | Negative(a) |                                 |                                                                                           |
|                               | Ames assay                           | S. Typhimurium TA98; TA100 | Not reported                    | Negative(c) |                                 | Kim et al. (1989)                                                                          |
|                               | Ames assay (preincubation method)    | S. Typhimurium TA98; TA100 | 0.4 μmol/plate (52 μg/plate)   | Negative(a) | Sasagawa and Matsushima (1991)  |                                                                                           |
|                               |                                      | S. Typhimurium TA100 | 100 μg/plate                    | Negative(b) | Ochiai et al. (1986)            |                                                                                           |
|                               |                                      | S. Typhimurium TA100 | Up to 3.33 mM (437 μg/mL)       | Negative(c) | Reddy et al. (2002)             |                                                                                           |
|                               | Mutation assay (paper-disk method)   | E. coli Sd-4-73     | 0.025 mL/disk                   | Negative   | Szybalski (1958)                |                                                                                           |
|                               | Chromosomal aberration assay         | Chinese hamster ovary cells | 1.3, 1.4 and 1.5 mM (+ S9) 1.4, 1.5 and 1.6 mM (− S9) | Positive(a) | Reddy et al. (2002)             | Aberrations were only detected at cytotoxic concentrations that showed marked inhibition of DNA synthesis |
|                               | Alkaline elution assay               | Rat primary hepatocytes (uninduced and PB/NF induced) | 0.5, 0.6, 0.7, 0.8, 0.9 and 1 mM | Negative   | Reddy et al. (2002)             | The study is considered valid                                                                |
|                               | DNA modification assay               | Isolated human genomic DNA | 25 and 500 μM (66 μg/mL)       | Positive(c) | Laws et al. (2001)              | Assay demonstrating inhibition of PCR amplification and spots demonstrated by postlabeling. Limited predictive value |
|                               | DNA Single-strand break assay        | Bovine kidney cells  | 10 μM–1 mM (131.2 μg/mL)        | Positive   | Kim et al. (1989)               | Abstract only. Validity cannot be evaluated                                                 |
| Chemical Name       | Test System                        | Test Object     | Concentration          | Result       | Reference                        | Comments                                                                 |
|---------------------|------------------------------------|-----------------|------------------------|--------------|-----------------------------------|--------------------------------------------------------------------------|
| (2-Acetylpyrrole    | Ames assay (plate incorporation    | S. Typhimurium  | 4, 20 and 100 µmol/plate| Negative     | Lee et al. (1994)                | Positive without S9 only at the two highest concentrations. High         |
| [14.047])           | method)                            | TA98            | (10913 µg/plate)       | Positive(c)  |                                   | concentrations. Technically acceptable, but of limited relevance due to high |
|                     |                                    | S. Typhimurium  | 100 µmol/plate (10,913 | Negative(a)  |                                   | concentrations.                                                          |
|                     |                                    | TA100           | µg/plate)              |              |                                   |                                                                          |
|                     | Ames assay                         | S. Typhimurium  | Up to 200 µg/plate     | Negative     | Wang et al. (1994)               |                                                                          |
|                     |                                    | TA98; TA100     |                        |(a)           |                                   |                                                                          |
| 2-Methylpyridine    | Ames assay (preincubation method)  | S. Typhimurium  | 3 µmol/plate (279 µg/plate) | Negative(a)  | Florin et al. (1980)          | Single-dose study                                                       |
| [14.134]            |                                    | TA98; TA100; TA1535; TA1537 |                      |              |                                   |                                                                          |
|                     | Ames assay (modified preincubation | S. Typhimurium  | 10 nmol-1 mmol/plate (| Negative(a)  | Aeschbacher et al. (1989) | The study is considered valid                                           |
|                     | method)                            | TA98; TA100; TA102 | 6 doses) (93 µg/mL) |              |                                   |                                                                          |
|                     | Ames assay (plate incorporation    | S. Typhimurium  | Up to 5,000 µg/plate (| Negative(a)  | Claxton et al. (1987)          | Individual dose levels not reported. The study is considered valid     |
|                     | method)                            | TA97; TA98; TA100; TA102 | 6 doses)              |              |                                   |                                                                          |
|                     |                                    | S. Typhimurium  | 50, 160, 500, 1,600 and | Negative(a)  | Vleminckx et al. (1993a)      | The study is considered valid                                           |
|                     |                                    | TA98; TA100; TA1535; TA1537 | 5,000 nL/plate (4,722 µg/mL) |              |                                   |                                                                          |
|                     | Mitotic aneuploidy assay           | S. cerevisiae   | 0.5–0.74% (6 doses) (6988 µg/mL) | Positive     | Zimmermann et al. (1986)       | Very high doses. The effect is considered thresholded. Limited          |
|                     | D61.M                              |                 |                        |              |                                   | relevance                                                               |
| HGPRT Gene mutation | Chinese hamster V79 lung cells     |                 | 4.5, 4.75, 5, 5.25 and | Negative(b)  | Vleminckx et al. (1993b)       | The study is considered valid                                           |
| assay               |                                    |                 | 5.5 µL/mL (5,194 µg/mL) |              |                                   |                                                                          |
| Alkaline elution    | Chinese hamster V79 lung cells     |                 | 2, 3, 4, 5 and 6 µL/mL (5,666 µg/mL) | Negative(b)  | Schriewer et al. (1993)      | The study is considered valid                                           |
| assay               |                                    |                 |                        |              |                                   |                                                                          |
| Chemical Name [FL-no] | Test System | Test Object | Concentration | Result | Reference | Comments |
|-----------------------|-------------|-------------|---------------|--------|-----------|----------|
| 3-Methylpyridine [14.135] | Ames assay (modified preincubation method) | *S. Typhimurium* TA98; TA100; TA1535; TA1537 | 85, 280, 840 and 8,540 µg/plate | Negative | Haworth et al. (1983) | The study is considered valid |
|                        | Ames assay (plate incorporation method) | *S. Typhimurium* TA98; TA100; TA1535; TA1537 | 50, 160, 500, 1,600 and 5,000 nL/plate (4,785 µg/plate) | Negative<sup>a</sup> | Vleminckx et al. (1993a) | The study is considered valid |
| 3-Ethylpyridine [14.061] | Mutagenicity assay | *E. coli* WP2 uvrA | 5-10 mg/plate (5000-10,000 µg/plate) | Negative | Pai et al. (1978) | Single-dose study. Very few experimental details. The validity cannot be evaluated |
|                        | HGPRT Gene mutation assay | Chinese hamster V79 lung cells | 3, 3.25, 3.5, 3.75 and 4 µL/mL (3,828 µg/mL) | Negative<sup>b</sup> | Vleminckx et al. (1993b) | The study is considered valid |
|                        | Alkaline elution assay | Chinese hamster V79 lung cells | 2, 3, 4 and 5 µL/mL (4,785 µg/mL) | Negative<sup>b</sup> | Schriewer et al. (1993) | The study is considered valid. |
| 4-Methylpyridine [14.136] | Ames assay (plate incorporation method) | *S. Typhimurium* TA98; TA100; TA1535; TA1537 | 50, 160, 500, 1,600 and 5,000 nL/plate (4779 µg/plate) | Negative<sup>a</sup> | Vleminckx et al. (1993a) | The study is considered valid. |
|                        | HGPRT Gene mutation assay | Chinese hamster V79 lung cells | 3.75, 4, 4.25 and 4.5 µL/mL (4,301 µg/mL) | Negative<sup>b</sup> | Vleminckx et al. (1993b) | The study is considered valid. |
|                        | Alkaline elution assay | Chinese hamster V79 lung cells | 3.75, 4, 4.25 and 4.5 µL/mL (4,301 µg/mL) | Negative<sup>b</sup> | Schriewer et al. (1993) | The study is considered valid. |
| (3-Ethylpyridine [14.061]) | Ames assay (preincubation method) | *S. Typhimurium* TA98; TA100; TA1535; TA1537 | 3 µmol/plate (321 µg/plate) | Negative<sup>a</sup> | Florin et al. (1980) | Single-dose study. |
| 2,4-Dimethylpyridine [14.104] | Mitotic aneuploidy assay | *S. cerevisiae* D61.M | 0.4–0.60% (6 doses) (5,551 µg/mL) | Positive | Zimmermann et al. (1986) | Very high doses. The effect is considered thresholded. Limited relevance |
| (2,6-Dimethylpyridine [14.065]) | Mitotic aneuploidy assay | *S. cerevisiae* D61.M | 0.5–0.60% (4 doses) (5,551 µg/mL) | Positive | Zimmermann et al. (1986) | Very high doses. The effect is considered thresholded. Limited relevance. |
| Chemical Name       | Test System                          | Test Object                                  | Concentration                  | Result           | Reference                        | Comments                                                                 |
|---------------------|--------------------------------------|----------------------------------------------|--------------------------------|------------------|-----------------------------------|--------------------------------------------------------------------------|
| 3,5-Dimethylpyridine | Ames assay (preincubation method)     | S. Typhimurium TA98; TA100; TA1535; TA1537   | 3 µmol/plate (321 µg/plate)   | Negative(a)      | Florin et al. (1980)              | Single-dose study                                                        |
|                     | Ames assay (plate incorporation method) | S. Typhimurium TA98; TA100; TA1535; TA1537; TA1538 | 100–10,000 µg/plate           | Negative         | Longfellow (1997)                | Very short summary. The results cannot be validated. High doses.         |
|                     | Mouse lymphoma assay                  | Mouse lymphocytes L5178Y tk<sup>+</sup>/tk<sup>−</sup> | 2,500–4,500 µg/mL (– S9) 1,000–4,000 µg/mL (+ S9) | Positive<sup>(a)</sup> |                                   | Very short summary. The results cannot be validated                     |
|                     | Mitotic aneuploidy assay              | S. cerevisiae D61.M                          | 0.5–0.87% (4 doses) (9,396 µg/mL) | Positive         | Zimmermann et al. (1986)         | Very high doses. The effect is considered thresholded. Limited relevance |
| (2-Acetylpyridine   | Mouse lymphoma assay                  | S. Typhimurium TA98; TA100; TA1535; TA1537; TA1538 | 100–10,000 µg/plate           | Negative         | Longfellow (1997)                | Very short summary. The results cannot be validated                     |
| [14.038])           | Mitotic aneuploidy assay              | S. cerevisiae D61.M                          | 0.5–1.11% (5 doses) (1,223 µg/mL) | Positive         | Zimmermann et al. (1986)         | Very high doses. The effect is considered thresholded. Limited relevance |
|                     | Mutation                              | E. coli WP2uvrA                               | 10,000 µg/plate               | Negative         | Pai et al. (1978)                | Single-dose study. Very few experimental details. The validity cannot be evaluated |
| (3-Acetylpyridine   | Mitotic aneuploidy assay              | S. cerevisiae D61.M                          | 0.5–1.11% (5 doses) (1,223 µg/mL) | Positive         | Zimmermann et al. (1986)         | Very high doses. The effect is considered thresholded. Limited relevance |
| [14.039])           | Mitotic aneuploidy assay              | S. cerevisiae D61.M                          | 0.5–1.11% (5 doses) (1,223 µg/mL) | Positive         | Zimmermann et al. (1986)         | Very high doses. The effect is considered thresholded. Limited relevance |
| 4-Acetylpyridine    | Ames assay (preincubation method)     | S. Typhimurium TA97; TA98; TA100; TA102; TA104; TA1535; TA1537; TA1538 | 5, 100, 300, 100, 3,000 and 10,000 µg/plate | Negative<sup>(a)</sup> | Zeiger et al. (1992)             | The study is considered valid                                            |
| [14.089]            | Mitotic aneuploidy assay              | S. cerevisiae D61.M                          | 0.5–1.11% (5 doses) (13,114 µg/mL) | Positive         | Zimmermann et al. (1986)         | Very high doses. The effect is considered thresholded. Limited relevance |
|                     | Mitotic aneuploidy assay              | S. cerevisiae D61.M                          | Up to 11 mg/mL                | Positive         | Whittaker et al. (1989)           | Purity 88%. Very high doses. The effect is considered thresholded. Limited relevance |

*: Supporting substances are listed in brackets.
(a): With and without metabolic activation.
(b): Without metabolic activation.
(c): With metabolic activation.
### Table A.3: Genotoxicity data (in vivo) EFSA/FGE.24Rev2 (EFSA CEF Panel, 2013)

| Chemical Name [FL-no]* | Test System | Test Object | Route | Dose | Result | Reference | Comments |
|------------------------|-------------|-------------|-------|------|--------|-----------|----------|
| (3-Methylindole [14.004])* | Micronucleus test | Mouse | Oral | 1,000 mg/kg day | Negative | (Reddy et al. (2003) Abstract only. The validity cannot be evaluated) |

*: Supporting substance.

### Table A.4: Genotoxicity data (in vitro) on 6-Methylquinoline

| Chemical Name [FL-no] | Test System | Test Object | Concentration | Result | Reference | Comments |
|-----------------------|-------------|-------------|----------------|--------|-----------|----------|
| 6-Methylquinoline [14.042] | Chromosomal aberration assay | Chinese hamster ovary cells | 52.7, 69.9, 174.8 and 349.5 μg/mL; 50.3, 125.5, 250.9 and 374.5 μg/mL | Negative (− S9) Positive (+ S9) | NTP (1986) |
| Sister chromatid exchanges | Chinese hamster ovary cells | 16.6, 25.1, 33 and 50 μg/mL; 16.7, 50.1, 166.9 and 500.7 μg/mL | Positive (− S9) Positive (+ S9) | NTP (1986) |
| Bacterial reverse mutation assay | S. Typhimurium TA98 and TA100 | 3, 10, 33, 100, 166, 333, 666, 1,000 μg/plate | Negative (− S9) Positive (+ S9) | NTP (1986) |

### Table A.5: Genotoxicity data (in vivo) on 6-Methylquinoline and 6-Isopropylquinoline

| Chemical Name [FL-no] | Test System | Test Object | Route | Dose | Result | Reference | Comments |
|-----------------------|-------------|-------------|-------|------|--------|-----------|----------|
| 6-Methylquinoline [14.042] | Micronucleus test | Male mice | Gavage | 0, 225, 450 and 900 mg/kg bw | Negative | Nakajima (2005) | Limited relevance. |
| Micronucleus test | NMRI mice | i.p. | 0, 286, 429, or 572 mg/kg bw | Negative | Wild et al. (1983) | Single intraperitoneal administration. Limited relevance |
| Comet assay in liver and duodenum | Male rat | Gavage | 100, 200 and 400 mg/kg bw | Equivocal in liver and Negative in duodenum | Keig-Shevlin (2016) | The study complies with OECD test guideline 489 and GLP guidelines |
| Gene mutation in liver and duodenum | Male Muta™ mice | Gavage | 125, 250, 500 mg/kg bw per day | Negative | Ballantyne (2017) | The study complies with OECD test guideline 488 and GLP guidelines |
| Micronucleus in peripheral blood reticulocytes | Male Muta™ mice | Gavage | 125, 250, 500 mg/kg bw per day | Negative |
| 6-Isopropylquinoline | Micronucleus test | NMRI mice | Oral | 0, 500, 1,000 and 2,000 mg/kg bw | Negative | Honarvar (2004) | Limited relevance |


### Appendix B – Summary of toxicity data

**Table B.1: Toxicity data considered by the panel**

| Chemical Name [FL-no] | Species; Sex No/group | Route | Doses (mg/kg bw per day) | Duration (days) | NOAEL (mg/kg bw per day) | Reference | Comments |
|-----------------------|------------------------|-------|--------------------------|-----------------|--------------------------|-----------|----------|
| Isoquinoline [FL-no: 14.001] | Rat; M, F 10 | Oral | 0, 0.03, 0.3 and 3 | 90 | 3 | Kojima (2006) | It is a GLP study performed according to the Japanese Guidelines for designation of food additives and revision of standards for use of food additives, Notification No 29 of the Environmental Health Bureau, Ministry of Health and Welfare, Japan, 22 March 1996. The requirements of this guideline are very similar to the OECD Guideline 408 |
| Pyrrole [FL-no: 14.041] | Rat; M, F 10 | Gavage | 0, 0.03, 0.3 and 3 | 90 | 3 | Marumo (2008) | Same as above |
| 2-Acetylpyrrole [FL-no: 14.047] | Rat; M, F 3 | Diet | 0, 85, 550 and 842 (males) 0, 91, 582 and 949 (females) | 14 | Range finding | Bauter (2012a) |
| | | Diet | 0, 68, 133 and 263 (males) 0, 79, 155 and 298 (females) | 90 | 48 | Bauter (2012b) | Compliant to the OECD test guideline 408 |
| 1-Furfurylpyrrole [FL-no: 13.134] | Rat; M, F 10 | Diet | 0, 29, 84 and 211 (males) 0, 27, 81 and 192 (females) | 14 | Range finding | Kappeler (2013a) | Compliant to the OECD test guideline 408 |
| | | Diet | 0, 25, 77 and 154 (males) 0, 25, 75 and 151 (females) | 90 | 25 | Kappeler (2013b) | |
| 6-methylquinoline [FL-no: 14.042] | Rat; M,F 10 to 16 | Diet | 2.2 (males) 2.7 (females) | 90 | 2.2 | Posternak et al. (1969) | The study was performed with a single dose that produced no adverse effects. The report is only a summary of the study. Poor study protocol |
| | | Diet | 25 mg/kg bw per day | 2-year | | Fukushima et al. (1981) | Carcinogenicity study. Only one dose was tested |
| | | Skin painting | 7.5 mg per animal | 18 weeks | | La Voie et al. (1984) | Assay for tumour-initiating activity |

M: Male.  
F: Female.
Table C.1: Summary of Safety Evaluation by JECFA (JECFA, 2005b)

| FL-no JECFA-no | EU Register name | Structural formula | EU MSDI (a) US MSDI (μg/capita per day) | Class (b) Evaluation procedure path (c) | Outcome on the named compound [(d) or (e)] | EFSA conclusion on the material of commerce |
|----------------|------------------|--------------------|----------------------------------------|----------------------------------------|---------------------------------------------|---------------------------------------------|
| 14.004 1304    | 3-Methylindole   | ![3-Methylindole](image) | 2.4 0.07 | Class I A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.007 1301    | Indole           | ![Indole](image) | 26 10  | Class I A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.041 1314    | Pyrrole          | ![Pyrrole](image) | 0.11 0.01 | Class I A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.038 1309    | 2-Acetylpyridine | ![2-Acetylpyridine](image) | 50 68  | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.039 1316    | 3-Acetylpyridine | ![3-Acetylpyridine](image) | 23 0.8  | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.045 1305    | 2-Acetyl-1-ethylpyrrole | ![2-Acetyl-1-ethylpyrrole](image) | 0.12 0.009 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.046 1306    | 2-Acetyl-1-methylpyrrole | ![2-Acetyl-1-methylpyrrole](image) | 1.2 0.02 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.047 1307    | 2-Acetylpyrrole  | ![2-Acetylpyrrole](image) | 3.3 0.2  | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.059 1312    | 3-Isobutylpyridine | ![3-Isobutylpyridine](image) | 0.049 0.07 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach |
| FL-no | JECFA-no | EU Register name | Structural formula | EU MSDI<sup>(a)</sup> (US MSDI<sup>(b)</sup> (µg/capita per day)) | Class<sup>(c)</sup> Evaluation procedure | Outcome on the named compound ['(d) or (e)'] | EFSA conclusion on the named compound (Procedure steps, intake estimates, NOAEL, genotoxicity) | EFSA conclusion on the material of commerce |
|-------|---------|------------------|-------------------|---------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|
| 14.060 | 1313    | 2-Pentylpyridine | ![structure](https://via.placeholder.com/150) | 0.061 0.07 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.061 | 1315    | 3-Ethylpyridine  | ![structure](https://via.placeholder.com/150) | 9.3 3 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.065 | 1317    | 2,6-Dimethylpyridine | ![structure](https://via.placeholder.com/150) | 0.26 0.007 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.066 | 1318    | 5-Ethyl-2-methylpyridine | ![structure](https://via.placeholder.com/150) | 0.12 0.04 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.068 | 1319    | 2-Propionylpyrrole | ![structure](https://via.placeholder.com/150) | 0.012 2 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.071 | 1320    | Methyl nicotinate | ![structure](https://via.placeholder.com/150) | 0.49 0.2 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.164 | 1322    | 2-Propylpyridine | ![structure](https://via.placeholder.com/150) | 0.61 0.9 | Class II A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.001 | 1303    | Isoquinoline | ![structure](https://via.placeholder.com/150) | 0.012 0.07 | Class III A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.042 | 1302    | 6-Methylquinoline | ![structure](https://via.placeholder.com/150) | 0.32 0.01 | Class III A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.058 | 1311    | 2-Isobutylpyridine | ![structure](https://via.placeholder.com/150) | 0.0061 0.9 | Class III A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| FL-no | JECFA-no | EU Register name | Structural formula | EU MSDI<sup>(a)</sup> (US MSDI) | Class<sup>(b)</sup> | Evaluation procedure path<sup>(c)</sup> | Outcome on the named compound<sup>(d,e)</sup> | EFSA conclusion on the named compound (Procedure steps, intake estimates, NOAEL, genotoxicity) | EFSA conclusion on the material of commerce |
|-------|---------|-----------------|-------------------|-----------------------------|----------------|-----------------------------------|---------------------------------|--------------------------------------------------------------------------------|---------------------------------|
| 14.072| 1321    | 2-(3-Phenylpropyl) pyridine | ![Structure](image) | 1.8 0.7 | Class III | A3: Intake below threshold | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 13.134| 1310    | 1-Furfurylpyrrole | ![Structure](image) | 0.12 0.07 | Class III | B3: Intake below threshold, B4: Adequate NOAEL exists | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |
| 14.030| 1308    | 2-Pyridine methanethiol | ![Structure](image) | 0.0012 0.007 | Class III | B3: Intake below threshold, B4: Adequate NOAEL exists | d | No safety concern at the estimated level of intake based on the MSDI approach | No safety concern at the estimated level of intake based on the MSDI approach |

<sup>(a)</sup>: EU MSDI: Amount added to food as flavour in (kg/year) × 10E9/(0.1 × population in Europe (≈ 375 × 10E6) × 0.6 × 365) = μg/capita per day. EU MSDIs may deviate from those reported in the JECFA evaluation because for several substances new data were available.

<sup>(b)</sup>: Thresholds of concern: Class I = 1,800 μg/person per day, Class II = 540 μg/person per day, Class III = 90 μg/person per day.

<sup>(c)</sup>: Procedure path A substances can be predicted to be metabolised to innocuous products. Procedure path B substances cannot.

<sup>(d)</sup>: No safety concern based on intake calculated by the MSDI approach of the named compound.

<sup>(e)</sup>: Data must be available on the substance or closely related substances to perform a safety evaluation.
Table C.2: Summary of safety evaluation by the EFSA (FGE.24Rev2) (EFSA CEF Panel, 2013)

| FL-no  | EU Register name           | Structural formula | MSDI<sup>(a)</sup> (µg/capita per day) | Class<sup>(b)</sup> Evaluation procedure path<sup>(c)</sup> | Outcome on the named compound [<sup>(d)</sup> or<sup>(e)</sup>] | Outcome on the material of commerce [<sup>(f)</sup>, <sup>(g)</sup>, or<sup>(h)</sup>] | Evaluation remarks |
|-------|---------------------------|--------------------|----------------------------------------|------------------------------------------------------------|---------------------------------------------------------------|---------------------------------------------------------------|-------------------|
| 14.110| Ethyl nicotinate          | ![Structure](structure1.png) | 0.013                                   | Class II A3: Intake below threshold                        | d                                                             | f                                                             |                   |
| 14.120| Isopropyl nicotinate      | ![Structure](structure2.png) | 0.0012                                  | Class II A3: Intake below threshold                        | d                                                             | f                                                             |                   |
| 14.023| 1-Methylpyrrole           | ![Structure](structure3.png) | 0.3                                     | Class II B3: Intake below threshold, B4: No adequate NOAEL | Additional data required                                      | i                                                             |                   |
| 14.085| 2-Acetyl-5-methylpyrrole  | ![Structure](structure4.png) | 0.0012                                  | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                                             | f                                                             |                   |
| 14.089| 4-Acetylpypidine          | ![Structure](structure5.png) | 0.0073                                  | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                                             | f                                                             |                   |
| 14.092| 2-Butylpyridine           | ![Structure](structure6.png) | 0.012                                   | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                                             | f                                                             |                   |
| 14.093| 3-Butylpyridine           | ![Structure](structure7.png) | 0.061                                   | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                                             | f                                                             |                   |
| 14.103| 2,3-Dimethylpyridine      | ![Structure](structure8.png) | 0.037                                   | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                                             | f                                                             |                   |
| 14.104| 2,4-Dimethylpyridine      | ![Structure](structure9.png) | 0.024                                   | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                                             | f                                                             |                   |
| 14.105| 3,4-Dimethylpyridine      | ![Structure](structure10.png) | 0.13                                    | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                                             | f                                                             |                   |
| FL-no  | EU Register name     | Structural formula | MSDI<sup>(a)</sup> (μg/capita per day) | Class<sup>(b)</sup> Evaluation procedure path<sup>(c)</sup> | Outcome on the named compound [(d) or (e)] | Outcome on the material of commerce [(f), (g), or (h)] | Evaluation remarks |
|--------|----------------------|--------------------|--------------------------------------|-------------------------------------------------|------------------------------------------|-------------------------------------------------|------------------|
| 14.106 | 3,5-Dimethylpyridine | ![Structure](image)  | 0.073                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.107 | 2,5-Dimethylpyrrole  | ![Structure](image) | 0.061                                | Class II B3: Intake below threshold, B4: No adequate NOAEL |                                           | Additional data required | i                |
| 14.115 | 2-Ethylpyridine      | ![Structure](image) | 0.027                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.116 | 4-Ethylpyridine      | ![Structure](image) | 0.027                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.117 | 2-Hexylpyridine      | ![Structure](image) | 0.012                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.118 | 2-Hydroxypyridine    | ![Structure](image) | 0.024                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.124 | 2-Isopropylpyridine  | ![Structure](image) | 0.021                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.125 | 4-Isopropylpyridine  | ![Structure](image) | 0.012                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.134 | 2-Methylpyridine     | ![Structure](image) | 0.21                                 | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.135 | 3-Methylpyridine     | ![Structure](image) | 0.027                                | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| 14.136 | 4-Methylpyridine     | ![Structure](image) | 0.73                                 | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                         | f                                               |                  |
| FL-no  | EU Register name                  | Structural formula | MSDI<sup>a</sup> (µg/capita per day) | Class<sup>b</sup> Evaluation procedure path<sup>c</sup> | Outcome on the named compound [d]<sup>(d)</sup> or [e]<sup>(e)</sup> | Outcome on the material of commerce [f]<sup>(f)</sup>, [g]<sup>(g)</sup>, or [h]<sup>(h)</sup> | Evaluation remarks |
|--------|-----------------------------------|--------------------|-------------------------------------|-------------------------------------------------|------------------------------------------------|------------------------------------------------|------------------|
| 14.140 | 3-Pentylpyridine                  | ![Structural formula](image1) | 0.0012                              | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                              | f                                              |                  |
| 14.143 | 3-Propylpyridine                  | ![Structural formula](image2) | 0.0012                              | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                              | f                                              |                  |
| 14.145 | Pyrrole-2-carbaldehyde            | ![Structural formula](image3) | 0.12                                | Class II B3: Intake below threshold, B4: No adequate NOAEL | Additional data required                      | j                                              |                  |
| 14.150 | 2,4,6-Trimethylpyridine           | ![Structural formula](image4) | 0.012                               | Class II B3: Intake below threshold, B4: Adequate NOAEL exists | d                                              | f                                              |                  |
| 14.169 | 1-Ethyl-2-pyrrolocarboxaldehyde   | ![Structural formula](image5) | 0.12                                | Class II B3: Intake below threshold, B4: No adequate NOAEL | Additional data required                      | j                                              |                  |
| 13.100 | 2-Acetyl-1-furfurylpyrrole        | ![Structural formula](image6) | 0.091                               | Class III B3: Intake below threshold, B4: No adequate NOAEL | Additional data required                      | i                                              |                  |
| 14.088 | 1-Acetylindole                    | ![Structural formula](image7) | 0.0012                              | Class III B3: Intake below threshold, B4: Adequate NOAEL exists | d                                              | f                                              |                  |
| 14.131 | 2-Methylindole                    | ![Structural formula](image8) | 0.0012                              | Class III B3: Intake below threshold, B4: Adequate NOAEL exists | d                                              | f                                              |                  |
| 14.163 | 1-Methylpyrrole-2-carboxaldehyde | ![Structural formula](image9) | 0.0024                              | Class III B3: Intake below threshold, B4: No adequate NOAEL | Additional data required                      | j                                              |                  |
| 14.002 | 4-Methylquinoline                 | ![Structural formula](image10) | 0.12                                | Class III No evaluation                          | i                                              | i                                              |                  |
| FL-no | EU Register name | Structural formula | MSDI<sup>(a)</sup> (µg/capita per day) | Class<sup>(b)</sup> Evaluation procedure path<sup>(c)</sup> | Outcome on the named compound [(d) or (e)] | Outcome on the material of commerce [(f), (g), or (h)] | Evaluation remarks |
|-------|----------------|--------------------|-------------------------------------|----------------------------------------|---------------------------------|-----------------------------------------------|-------------------|
| 14.094 | 4-Butylquinoline | ![4-Butylquinoline](image) | 0.0012 | Class III No evaluation | | | i |
| 14.138 | 2-Methylquinoline | ![2-Methylquinoline](image) | 0.012 | Class III No evaluation | | | i |

(a): EU MSDI: Amount added to food as flavour in (kg/year) × 10E9/(0.1 × population in Europe (= 375 × 10E6) × 0.6 × 365) = µg/capita per day.
(b): Thresholds of concern: Class I = 1,800 µg/person per day, Class II = 540 µg/person per day, Class III = 90 µg/person per day.
(c): Procedure path A substances can be predicted to be metabolised to innocuous products. Procedure path B substances cannot.
(d): No safety concern based on intake calculated by the MSDI approach of the named compound.
(e): Data must be available on the substance or closely related substances to perform a safety evaluation.
(f): No safety concern at the estimated level of intake of the material of commerce meeting the specification requirement (based on intake calculated by the MSDI approach).
(g): Tentatively regarded as presenting no safety concern (based on intake calculated by the MSDI approach) pending further information on the purity of the material of commerce and/or information on stereoisomerism.
(h): No conclusion can be drawn due to lack of information on the purity of the material of commerce.
(i): No longer supported by Industry (DG SANCO, 2012).
(j): No longer supported by Industry (DG SANCO, 2013).
### Appendix D – Exposure data

**Table D.1**: Normal and Maximum use levels available for substances in FGE.77Rev3

| FL-no | Normal use levels (mg/kg) | Maximum use levels (mg/kg) |
|-------|---------------------------|----------------------------|
|       | 01.0 | 02.0 | 03.0 | 04.1 | 04.2 | 05.0 | 06.0 | 07.0 | 08.0 | 09.0 | 10.0 | 11.0 | 12.0 | 13.0 | 14.1 | 14.2 | 15.0 | 16.0 |
| 13.134| 0.011| 0.13 | 0.27 |     |     |     |     | 18  |     |     |     |     |     |     |     |     |     |     |     |
|       | 0.027| 0.13 | 0.27 |     |     |     |     | 880 |     |     |     |     |     |     |     |     |     |     |     |
| 14.001|     | 0.025|     |     |     |     |     |     | 0.25|     |     |     |     |     |     |     |     |     |     |
|       |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| 14.030|     | 1    |     |     |     |     |     | 1   |     |     |     |     |     |     |     |     |     |     |     |
|       |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| 14.041|     |     |     |     |     |     |     | 1   |     |     |     |     |     |     |     |     |     |     |     |
|       |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| 14.042|     |     |     |     |     |     |     | 1.2 |     |     |     |     |     |     |     |     |     |     |     |
|       |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| 14.058|     |     |     |     |     |     |     | 0.025|     |     |     |     |     |     |     |     |     |     |     |
|       |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| 14.072|     |     |     |     |     |     |     | 0.2 |     |     |     |     |     |     |     |     |     |     |     |
Table D.2: Estimated intakes based on the MSDI and the mTAMDI\(^{(a)}\) approach – FGE.77Rev3

| FL-no | EU Register name                  | MSDI – EU (µg/capita per day) | MSDI – USA (µg/capita per day) | mTAMDI (µg/person per day) | Structural class | Threshold of concern (µg/person per day) |
|-------|-----------------------------------|--------------------------------|--------------------------------|---------------------------|-----------------|----------------------------------------|
| 14.004| 3-Methylindole                    | 2.4                            | 0.07                           |                           | Class I         | 1,800                                  |
| 14.007| Indole                            | 26                             | 10                             |                           | Class I         | 1,800                                  |
| 14.041| Pyrrole                           | 0.11                           | 0.01                           | 480                       | Class I         | 1,800                                  |
| 14.038| 2-Acetylpyridine                  | 50                             | 68                             |                           | Class II        | 540                                    |
| 14.039| 3-Acetylpyridine                  | 23                             | 0.8                            |                           | Class II        | 540                                    |
| 14.045| 2-Acetyl-1-ethylpyrrole           | 0.12                           | 0.009                          |                           | Class II        | 540                                    |
| 14.046| 2-Acetyl-1-methylpyrrole          | 1.2                            | 0.02                           |                           | Class II        | 540                                    |
| 14.047| 2-Acetylpyrrole                   | 3.3                            | 0.2                            |                           | Class II        | 540                                    |
| 14.059| 3-Isobutylpyridine                | 0.049                          | 0.07                           |                           | Class II        | 540                                    |
| 14.060| 2-Pentylypyridine                 | 0.061                          | 0.07                           |                           | Class II        | 540                                    |
| 14.061| 3-Ethylpyridine                   | 9.3                            | 3                              |                           | Class II        | 540                                    |
| 14.065| 2,6-Dimethylpyridine              | 0.26                           | 0.007                          |                           | Class II        | 540                                    |
| 14.066| 5-Ethyl-2-methylpyridine          | 0.12                           | 0.04                           |                           | Class II        | 540                                    |
| 14.068| 2-Propionylpyrrole                | 0.012                          | 0.2                            |                           | Class II        | 540                                    |
| 14.071| Methyl nicotinate                 | 0.49                           | 0.2                            |                           | Class II        | 540                                    |
| 14.164| 2-Propylpyridine                  | 0.61                           | 0.9                            |                           | Class II        | 540                                    |
| 14.001| Isoquinoline                      | 0.012                          | 0.07                           | 64                        | Class III       | 90                                     |
| 14.042| 6-Methylquinoline                 | 0.32                           | 0.01                           | 830                       | Class III       | 90                                     |
| 14.058| 2-Isobutyrylidine                 | 0.0061                         | 0.9                            | 64                        | Class III       | 90                                     |
| 14.072| 2-(3-Phenylpropyl)pyridine        | 1.8                            | 0.7                            | 65                        | Class III       | 90                                     |
| 14.134| 1-Furfurylpyrrole                 | 0.12                           | 0.07                           | 3000                      | Class III       | 90                                     |
| 14.030| 2-Pyridine methanethiol           | 0.0012                         | 0.007                          | 520                       | Class III       | 90                                     |

\(^{(a)}\): For the calculation of mTAMDI – see e.g. Annex II in FGE.03 (EFSA, 2004).