

UNIVERSITY OF BIRMINGHAM

Research at Birmingham

Emissions and indoor concentrations of particulate matter and its specific chemical components from cooking

Abdullahi, Lami; Delgado Saborit, Juana Maria; Harrison, Roy

DOI:

[10.1016/j.atmosenv.2013.01.061](https://doi.org/10.1016/j.atmosenv.2013.01.061)

License:

None: All rights reserved

Document Version

Peer reviewed version

Citation for published version (Harvard):

Abdullahi, L, Delgado Saborit, JM & Harrison, RM 2013, 'Emissions and indoor concentrations of particulate matter and its specific chemical components from cooking: A review', *Atmospheric Environment*, vol. 71, pp. 260-294. <https://doi.org/10.1016/j.atmosenv.2013.01.061>

[Link to publication on Research at Birmingham portal](#)

Publisher Rights Statement:

NOTICE: this is the author's version of a work that was accepted for publication in *Atmospheric Environment*. Changes resulting from the publishing process, such as peer review, editing, corrections, structural formatting, and other quality control mechanisms may not be reflected in this document. Changes may have been made to this work since it was submitted for publication. A definitive version was subsequently published in *Atmospheric Environment*, [vol. 71, 2013] DOI: <http://dx.doi.org/10.1016/j.atmosenv.2013.01.061>

General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact UBIRA@lists.bham.ac.uk providing details and we will remove access to the work immediately and investigate.

1
2
3
4
5
6
7
8

EMISSIONS AND INDOOR CONCENTRATIONS OF PARTICULATE MATTER AND ITS SPECIFIC CHEMICAL COMPONENTS FROM COOKING: A REVIEW

9
10
11
12

**Karimatu L. Abdullahi¹,
Juana Maria Delgado-Saborit¹
and Roy M. Harrison^{*1,2}**

13
14
15
16
17
18
19
20

**¹Division of Environmental Health & Risk Management
School of Geography, Earth & Environmental Sciences
University of Birmingham, Edgbaston
Birmingham B15 2TT
United Kingdom**

* To whom correspondence should be addressed

Tele: +44 121 414 3494; Fax: +44 121 414 3708; Email: r.m.harrison@bham.ac.uk; LKA010@bham.ac.uk;
j.m.delgadosaborit@bham.ac.uk

² Also at: Department of Environmental Sciences / Center of Excellence in Environmental Studies, King
Abdulaziz University, PO Box 80203, Jeddah, 21589, Saudi Arabia.

21

22 **ABSTRACT**

23 It has long been known that cooking can create high concentrations of aerosol indoors.
24 Increasingly, it is now being reported that cooking aerosol is also a significant component of
25 outdoor particulate matter. As yet, the health consequences are unquantified, but the
26 presence of well known chemical carcinogens is a clear indication that cooking aerosol
27 cannot be benign. This review is concerned with current knowledge of the mass
28 concentrations, size distribution and chemical composition of aerosol generated from typical
29 styles of cooking as reported in the literature. It is found that cooking can generate both
30 appreciable masses of aerosol at least within the area where the cooking takes place, that
31 particle sizes are largely within the respirable size range and that major groups of chemical
32 compounds which have been used to characterise cooking aerosol include alkanes, fatty
33 acids, dicarboxylic acids, lactones, polycyclic aromatic hydrocarbons, alkanones and sterols.
34 Measured data, cooking emission profiles and source apportionment methods are briefly
35 reviewed.

36

37 **Keywords:** Cooking; cooking aerosol; cooking emissions; cooking tracers

38

40 1. INTRODUCTION

41 Indoor air pollution is reported to be responsible for 2.7% of the global burden of disease
42 (Fang et al., 1999). Worldwide more than 1 million people die from chronic obstructive
43 pulmonary disease (COPD) annually due to indoor exposure to smoke which generally
44 contains a range of health-damaging pollutants, such as fine particles and carbon monoxide
45 (Hetland et al., 2000). The use of solid fuels (biomass and coal) for cooking and heating
46 homes is practised by around 3 billion people in open fires and leaky stoves, especially by
47 people with low and medium resources in developing countries. As such, poorly ventilated
48 homes can have indoor smoke concentrations of respirable particles of more than 100 times
49 the acceptable levels (Hetland et al., 2000) with mostly women and young children being
50 exposed to these extremely high levels.

51 Indoor levels of particles in developed countries are much lower than in developing countries
52 and this is generally attributable to the advancement in technology for general household
53 activities and also the use of cleaner fuels (such as liquefied petroleum gas, electricity and
54 natural gas) for cooking and heating. However, there are still observed risks to health in
55 people exposed to indoor air in these locations. Legislation relating to air pollutant exposure
56 in developed countries is normally based upon ambient outdoor concentrations, potentially
57 leading to inadequate protection of the general public who spend the majority of their time at
58 home, offices or other enclosed locations where the concentrations of some pollutants are
59 often much higher than ambient levels (Marcazzan et al., 2001). Knowledge of the indoor
60 environment is limited and is of great importance as the majority of people have been found
61 to spend about 80-90% of their time indoors in many countries (Koistinen K.J., 2001;
62 Scapellato et al., 2009; Delgado-Saborit et al., 2011). Also the indoor environments have

63 been found to be affected by factors such as the design of the buildings, insulation and
64 ventilation in order to ensure an adequately controlled environment for thermal comfort,
65 which can also affect level of individual exposure (Tan et al., 2012).

66 This literature review deliberately omits those studies that report emissions from cooking
67 using biomass and solid fuels. In such studies, the elevated concentrations reported derive
68 largely from the combustion of the fuel used for cooking rather than from the cooking itself.
69 This study focuses on studies of cooking emissions reported from the use of cleaner cooking
70 fuels, such as electricity and gas. It is expected that the use of electricity and gas will also
71 contribute to the cooking emissions. However, although such emissions will be included in
72 the concentrations reported in the literature, the main contribution to those concentrations is
73 expected to be from compounds deriving from the cooking of the ingredients itself.

74 **1.1. Particulate Matter**

75 Particulate matter (PM) is defined as the mass of a mixture of solid particles and liquid
76 droplets of various sizes (range from a few nanometres to tens of micrometres) suspended in
77 a volume of air which represent a broad class of chemically and physically diverse
78 substances. Particulate matter is classified according to its size, thus PM_{10} is defined as the
79 concentration of particulate matter with aerodynamic diameter of 10 micrometres or less,
80 while $PM_{2.5}$ is defined as the concentration of particulate matter that has aerodynamic
81 diameter of 2.5 micrometres or less.

82 Particulate matter consists of components that are released directly from a source (primary
83 PM) or are formed by chemical reactions in the atmosphere (secondary PM). It comes from
84 natural and man-made sources and consists of a range of chemical compounds which can be
85 useful for the identification of the source. Primary particulate matter is released from sources

86 which include road transport (tyres and brake wear, engine combustion, road dusts),
87 industrial, commercial and domestic burning of fuels and also dust from these activities, and
88 natural sources (sea spray and dust).

89 **1.2. Cooking and PM**

90 Several studies of indoor air have identified cooking as one of the most significant particle
91 generating activities indoors (Kamens et al., 1991; Ozkaynak et al., 1996; Chao and Cheng,
92 2002; Lazaridis et al., 2006; Zhao et al., 2007a; Buonanno et al., 2009; Lai et al., 2010; Wan
93 et al., 2011; Massey et al., 2012). The general population is exposed to cooking-related risk
94 regardless of race, age, wealth and cultural food preferences as cooking is an important aspect
95 of human culture (Kim et al., 2011). The processes used in cooking such as frying, roasting,
96 grilling, boiling and broiling contribute to pollutant emissions and are affected by ingredients,
97 recipes and procedures, fuel types, temperature and extraction/ventilation equipment (Zhang
98 et al., 2010). Table 1 summarises the cooking styles, ingredients and oils used for some
99 common cultural culinary techniques. The table shows that many of the ingredients and
100 cooking methods are common to various culinary techniques. This is a reflection of the
101 current intercultural exchange that represents an increasingly vague difference between the
102 major styles of cooking.

103 Cooking contributes particles to outdoor as well as indoor air. Commercial cooking
104 emissions may have contributed to the exceedence of the Federal PM_{2.5} air quality standards
105 in certain regions such as Pittsburgh, Pennsylvania, where meat charbroiling was shown to
106 contribute to carbonaceous PM (Cabada et al., 2002). Large scale cooking has been identified
107 to be an important contributor to organic carbon (OC) and elemental carbon (EC) in the urban
108 environment including secondary organic aerosol (SOA) formed by condensation of reaction
109 products of gaseous organic emissions following photochemical processes) (Roe et al., 2005).

110 Rogge et al. (1991) reported that 21% of the primary fine organic aerosol in the Los Angeles
111 area in the 1980s was generated by charbroiling and meat cooking activities, which was in
112 agreement with previous studies in the area (Hildemann et al., 1991b). A similar study in
113 1997 in Denver Colorado, the Northern Front Range Air Quality Study (NFRAQS), found
114 that meat cooking contributed about 15% of PM_{2.5} organic aerosol concentrations (Watson et
115 al., 1998).

116 Recently a study in New York City (NYC) using a High-Resolution Time-of-Flight Aerosol
117 Mass Spectrometer (HR-ToF-AMS) identified that cooking and traffic were two distinct and
118 mass-equivalent Primary Organic Aerosol sources, contributing 30% of the total Organic
119 Aerosol (OA) mass collectively during the period (Sun et al., 2011). The average mass
120 concentration of Cooking OA was 1.02 µg/m³, higher than the mass concentration of
121 Hydrocarbon like OA (0.91 µg/m³), which was surprising as the sampling site was actually
122 close to two major highways (<1 mile), giving a clear indication that cooking activities were
123 an important source of primary particles in NYC. An earlier air quality campaign in Beijing
124 in 2008 using a HR-ToF-AMS found that 24.4% of total organic mass was similarly
125 attributed to cooking related organic aerosols (Huang et al., 2010) .

126 Measurement of particle number and size distribution of particles generated during cooking
127 has been carried out in various studies to provide a better understanding of characteristics of
128 particles generated during cooking (Abt et al., 2000; Buonanno et al., 2009; Dennekamp et
129 al., 2001; See and Balasubramanian, 2006a; Wallace et al., 2004; He et al., 2004b).

130 Generally the risk associated with cooking is still poorly understood, although such
131 awareness is necessary to ensure adequate protection of health for the general public. This
132 review will provide an analysis of the sampling methods, data upon aerosols emitted from

133 cooking, the source profiles identified and their usefulness in apportionment studies will also
134 be considered.

135

136 **2. EMISSIONS FROM COOKING**

137 Studies of cooking emissions have been carried out in both real life kitchens and in controlled
138 environments. It is assumed that in controlled experimental setups, the measurements are
139 influenced mainly by the fuel used and the food being cooked while in actual real life
140 kitchens measurement of emissions are influenced by many factors such as room
141 arrangement, building materials, outdoor infiltration, other combustion devices, ventilation,
142 and cooking methods (Huboyo et al., 2011).

143 Visible fumes are generated during the cooking process, which are usually due to
144 submicrometer sized particles, which consist of oil droplets, combustion products, steam
145 from water in the food being cooked and condensed organic pollutants. The particulate matter
146 (PM) generated is generally within the ultrafine particle (UFP) - which represents particles of
147 diameter less than 100 nm - and fine PM (PM_{2.5}) size ranges. The physical stirring of food has
148 been found to lead to the generation of large aerosols due to the process of splashing of the
149 ingredients (Long et al., 2000). The combustion process associated with cooking can lead to
150 the formation and direct emission of ultrafine particles (UFP) to the atmosphere, and hot
151 vapours in the cooking fumes may also cool and nucleate to form more UFP (Sioutas et al.,
152 2005; Lai and Ho, 2008). These particles may contain organic substances, such as polycyclic
153 aromatic hydrocarbons (PAH) and heterocyclic amines, adsorbed on their surfaces (Ho et al.,
154 2002).

155 There is a scarcity of national inventories of cooking activities, but an attempt was made by
156 Roe et al. (2005) to compile a national emission inventory for commercial cooking in the
157 United States as listed in Table 2. For comparison, data for highway vehicles extracted from
158 the National Emissions Inventory (NEI) Air Pollutant Emissions Trends Data for the same
159 year (Chappell et al., 2003), show that although traffic emits orders of magnitude more CO
160 and VOC than cooking, particulate matter emissions from cooking are comparable with those
161 emitted from highway vehicles. This is consistent with a study of Li et al. (2003), who found
162 that the emission rates of total PAH from cooking sources in the study city (i.e. emissions
163 from both restaurants and home kitchens), were slightly lower than those for traffic sources in
164 a representative city of Taiwan (8,973 kg/year for cooking against 13,500 kg/year for traffic).
165 Nonetheless, they observed that the emission rate for B[a]P_{eq} toxic equivalent for cooking
166 sources was much higher than that from traffic sources (675 kg/year from cooking and 61.4
167 kg/year emitted from traffic sources). This indicated that cooking PAH may cause much more
168 serious problems than traffic sources in terms of carcinogenic potency (Li et al., 2003).

169

170 **2.1. Particle mass concentration**

171 The PTEAM Study (Particle Total Exposure Assessment Methodology) performed in the US,
172 reported around 20 $\mu\text{g}/\text{m}^3$ higher particle concentrations in houses where cooking took place
173 during their monitoring than those house where no cooking occurred (Wallace, 1989). They
174 reported that the proportion of PM_{2.5} and PM₁₀ due to cooking represented 25% for both
175 particle sizes. This proportion increased to 65% and 55%, respectively, when considering
176 indoor sources alone (Ozkaynak et al., 1996). Source apportionment of PTEAM ambient and
177 personal exposure samples using a combined receptor model found that cooking was the
178 largest contributory source of PM indoors, responsible for about 52.5% of the personal

179 exposure samples and 43.2% of residential indoor concentrations (Zhao et al., 2006). After
180 1,000 hours of cooking, they also found that the mean $PM_{2.5}$ personal exposure increased an
181 average of $56 \mu\text{g}/\text{m}^3$ while cooking activities took place, and that cooking increased the
182 overall 24-hours personal exposure about $2.5 \mu\text{g}/\text{m}^3$ in those persons that had cooked during
183 the sampling day (Wallace et al., 2006).

184 A study to characterize indoor sources of particles conducted in Boston, USA, made
185 measurements of particle size and volume concentration over 6 days in four non-smoking
186 households equipped with gas and electric stoves (Abt et al., 2000). The monitoring
187 equipment was placed in a single indoor location adjacent to the kitchen and living room and
188 from the data obtained, it was found that the highest mean peak mass concentrations were for
189 barbequing and sautéing for the $PM_{0.02-0.5}$ and $PM_{0.7-10}$ respectively, whilst the lower mean
190 peak concentrations were found for frying and oven cooking or toasting for the same size
191 ranges respectively (see Table 3) (Abt et al., 2000).

192 Another US study found that the average $PM_{2.5}$ concentration due to cooking over 195
193 cooking events was about $5.5 \mu\text{g}/\text{m}^3$ with a standard error of $2.3 \mu\text{g}/\text{m}^3$ (Allen et al., 2004). In
194 Europe, a study made a comparison of elderly residents in Amsterdam (47) and Helsinki (37),
195 and found that the estimated contribution from cooking ranged from $1.9 \mu\text{g}/\text{m}^3$ for indoor
196 $PM_{2.5}$ in Helsinki to $3.4 \mu\text{g}/\text{m}^3$ for $PM_{2.5}$ personal exposure concentrations (Brunekreef et al.,
197 2005).

198 Rates of emission of aerosol have been reported to vary based on type of appliance used, the
199 cooking conditions used and fat content of meat (McDonald et al., 2003). In an experiment
200 where hamburger, steak and chicken were grilled and charbroiled, McDonald et al. (2003)
201 found that the $PM_{2.5}$ emission rate for charbroiling meats ranged between 4.4 to 15 g/kg. The
202 largest quantity of $PM_{2.5}$ was emitted by hamburger (15 g/kg) which had higher fat content

203 (30%) and were cooked on a char broiler. These results are consistent with data reported by
204 Hildemann et al. (1991a) . McDonald et al. (2003) reported that charbroiling produced higher
205 concentrations than frying, 12-46 g/kg meat when charbroiling vs. 0.57 g/kg meat when
206 frying. They also reported that charbroiling lean meat produced less concentrations of
207 particles in the smaller size range (<20 nm) and in the larger size range (>100 nm) than
208 regular meat.

209 Similarly, Buonanno et al. (2009) found that gas stoves emitted more particles than an
210 electric stove when frying resulting in higher indoor concentrations when gas stoves were
211 used (60-118 $\mu\text{g}/\text{m}^3$) than when electric stoves were employed (12-27 $\mu\text{g}/\text{m}^3$); and that
212 emission rates were considerably affected by the type of food used (Table 3). Increased
213 emissions measured at the source were reported to be a function of increased cooking
214 temperature. Foods containing a higher percentage of fat generated higher emission rates than
215 those with less fat percentage. They reported higher aerosol mass emission when cooking
216 fatty foods resulting in higher indoor concentrations (280-389 $\mu\text{g}/\text{m}^3$) than when cooking
217 vegetables (78 $\mu\text{g}/\text{m}^3$). Particle emission factor varied significantly also with type of oil used.
218 Sunflower oil generated the lowest mass emission factors, whilst the highest emissions were
219 from olive oil (Buonanno et al., 2009). Glytsos (2010) reported that frying of onions in olive
220 oil in a controlled room emitted $\text{PM}_{2.5}$ increasing the indoor concentration in the range of 70
221 to 600 $\mu\text{g}/\text{m}^3$ (Glytsos et al., 2010).

222 See and Balasubramanian (2006b) investigated the physical and chemical properties of
223 emissions from a Chinese food stall in Singapore while food was stir fried in a wok using a
224 gas stove, and at two different and distinct times (See and Balasubramanian, 2006b). The
225 mass concentration of particles ($\text{PM}_{2.5}$) measured in the food stall at the opposite site of a 4-

226 LPG burner stove increased from 26.7 $\mu\text{g}/\text{m}^3$ during non-cooking hours to 312.4 $\mu\text{g}/\text{m}^3$ during
227 cooking hours (increased by a factor of 12).

228 Analysis of various cooking methods which included steaming, boiling, stir-frying, pan-
229 frying and deep-frying revealed that the largest amount of particulate matter measured at 20
230 cm from the cooker was generated during deep frying ($\text{PM}_{2.5}$ 190 $\mu\text{g}/\text{m}^3$) and the lowest
231 concentration was observed during steaming ($\text{PM}_{2.5}$ 72 $\mu\text{g}/\text{m}^3$) (See and Balasubramanian,
232 2008). Both studies have indicated that cooking with oil contributes to the production of more
233 particles than cooking with water, which is consistent with the work of He et al. (2004a). In
234 another study, See et al. (2006) made a comparison of emissions from Chinese, Indian and
235 Malay food stalls and reported that the highest mass concentrations of $\text{PM}_{2.5}$ were found in
236 the Malay stall (245.3 $\mu\text{g}/\text{m}^3$), whilst the lowest were measured in the Indian stall (186.9
237 $\mu\text{g}/\text{m}^3$) (See et al., 2006).

238 Several studies have found that Asian style cooking emits more particulate matter than
239 Western cooking with concentrations of $\text{PM}_{2.5}$ ranging 30 to 1,400 and 20 to 535 $\mu\text{g}/\text{m}^3$ as
240 reported by various groups (Lee et al., 2001; Levy et al., 2002; He et al., 2004b)

241 A summary of the main studies reporting aerosol concentration emitted from cooking and the
242 reported concentrations can be found in Table 3.

243

244 **2.2. Particle size distribution**

245 The size distribution of aerosols emitted from cooking activities has been reported in several
246 studies whose methodology and study description is summarised in Table 4 and results are
247 compiled in Table 5. Generally some of these studies have shown that indoor particle
248 concentrations are substantially affected by cooking activities, cleaning and the movement of

249 people (Abt et al., 2000; He et al., 2004a; Diapouli et al., 2011). The largest percentage of the
250 measured particles are ultrafine particles (UFP), with modes in the number distribution
251 reported generally in the range of 20 to 100 nm (see Table 5).

252 He et al. (2004a) studied 15 homes in Australia while cooking was carried out under good
253 and poor ventilation for 48 hours. They found that some indoor activities led to an increase in
254 indoor particle number concentration of about 1.5-27 times concentrations in comparison
255 with the particle number concentration when no indoor source was in operation. They also
256 found an emission rate ranging $0.2-4 \times 10^{12}$ particles/min and peak submicron number
257 concentrations for cooking of 16,000 and 180,000 part/cm³ (He et al., 2004a).

258 An investigation of the size distribution of particles emitted from cooking was carried out
259 using a scanning mobility particle sizer (SMPS) in a domestic kitchen using five different
260 cooking methods, such as steaming, boiling, stir-frying, pan-frying, and deep-frying. Deep-
261 frying was found to have the highest particle number concentration, whilst steaming
262 produced the lowest particle number concentration. Their observations found that cooking
263 activities using oil produce higher concentrations than those using water (See and
264 Balasubramanian, 2006a). They reported a 24-fold increase in particle concentration observed
265 between deep frying and background concentrations (6.0×10^5 cm⁻³ compared to background
266 concentrations which were 2.5×10^4 cm⁻³) (See and Balasubramanian, 2006a). In another
267 study, they characterised Chinese cooking emissions, and found that the average number
268 concentration increased by factor a of 85 during the cooking periods (7.7×10^5 part/cm³
269 compared to 9.1×10^3 part/cm³ during non- cooking hours (See and Balasubramanian, 2006b).

270 Yeung and To (2008) examined aerosols generated by commercial food preparation and
271 found a lognormal size distribution. Increased cooking temperature resulted in an increased
272 modal diameter of aerosols. Higher cooking temperature also increased the normalized

273 number concentration sub-micrometer aerosols (between 0.1 and 1.0 μm) (Yeung and To,
274 2008).

275 Siegmann and Sattler (1996) found that diameter and number concentrations of oil droplets
276 increased with an increase in temperature. They analysed aerosols from different hot
277 vegetable oils and obtained a size distribution with a mean droplet size range of 30 nm at
278 223°C to 100 nm at 256°C. Particle number concentration increased from $2.25 \times 10^5 \text{ part/cm}^3$
279 to $4.5 \times 10^5 \text{ part/cm}^3$ in the same range of temperatures (Siegmann and Sattler, 1996).

280 Dennekamp et al. (2001) studied the generation of ultrafine particles and nitrogen oxides
281 using different cooking procedures comparing gas and electric stoves in a laboratory. They
282 found higher concentrations of particles in the size range of 15-40nm (and also oxides of
283 nitrogen) when cooking on gas (Dennekamp et al., 2001). The smaller particles generated
284 were found to grow in size with time during the experiment. The high concentrations of
285 pollutants observed were attributed to the absence of ventilation in their laboratory kitchen.

286 Frying of onions in olive oil in a controlled room to characterise contributors of particle
287 concentrations in indoor environments produced high particle concentrations, ranging
288 between $9 - 15 \times 10^4 \text{ particles cm}^{-3}$ (Glytsos et al., 2010). High emission of nanoparticles
289 were reported during frying ($1.15 \times 10^5 \text{ part/cm}^3$, mainly 20 nm). However, sometime after
290 the frying stopped (i.e. 45 min later), the number concentration decreased down to 4×10^5
291 part/cm^3 and particles become larger leading to a bimodal size distribution indicating a strong
292 coagulation effect (Glytsos et al., 2010), which is consistent with previous studies
293 (Dennekamp et al., 2001; Sjaastad et al., 2008).

294 A study in an apartment in Taiwan found a range of mode diameters of particles
295 concentrations between 30-50nm for domestic cooking processes of scrambling eggs, frying
296 chicken, and cooking soup with higher mode diameter for frying chicken (Li et al., 1993).

297 Similarly, in an 18 month campaign in a four bedroom house consisting of three levels
298 located near Washington DC, USA; particles generated from cooking were found to be
299 mainly in the ultrafine range (about 90% of total particles), with frying being found to
300 generate more particles than any other cooking method (Wallace et al., 2004), consistent with
301 recent studies (Hussein et al., 2006; Buonanno et al., 2011; Huboyo et al., 2011).

302 Buonanno et al. (2009) sought to evaluate the influence of temperature, oil, food and stove
303 type on particle number, surface area and mass emission factors consequence of cooking with
304 different methods such as grilling and frying. They used a Scanning Mobility Particle Sizer
305 (SMPS) and Aerodynamic Particle Sizer (APS). They found that frying food with oil using an
306 electrical frying pan produced emission factors well below those observed for frying using a
307 gas stove. The particle emission factor was also dependent upon the temperature of the stove,
308 with values 9 and 4 times higher at the maximum stove power for gas and electric stoves
309 respectively (Buonanno et al., 2009), consistent with previous studies of Siegmann and
310 Sattler (1996), Dennekamp et al. (2001), Yeung and To (2008) and To and Yeung (2011). In
311 another study by them, they reported high particle indoor concentrations (3×10^4 – 6×10^5
312 particles cm^{-3}) in 14 pizzerias and PM_{10} concentrations of about 10–327 $\mu\text{g m}^{-3}$ during
313 normal ventilation conditions (Buonanno et al., 2010). However most of the particles
314 generated in this study are believed to be from the wood burning used to fire the oven, and
315 highlights the high particle concentrations that can build up in such microenvironments. In
316 another study, Buonanno and colleagues found that frying the same type of food consistently
317 emitted more particles than grilling, with a factor of 1.4-1.5 (Buonanno et al., 2011).

318

319

320 3. ON-LINE CHEMICAL CHARACTERISATION – AEROSOL MASS 321 SPECTROMETER

322 Atmospheric organic aerosol consists of many compounds with dissimilarities in properties
323 making the characterisation of its chemical composition and mass concentration by analytical
324 processes quite difficult (Seinfeld and Pankow, 2003; Kanakidou et al., 2005; Goldstein and
325 Galbally, 2007; Zhang et al., 2011). It has been found that only about 10% of the Organic
326 Aerosol (OA) mass can be easily speciated by techniques such as GC-MS and inconsistencies
327 have been found to exist between different thermal-optical organic carbon (OC)
328 quantification and artefact removal techniques (Schauer et al., 1996; Zhang et al., 2007). In
329 order to obtain better understanding of submicron aerosols, on-line chemical characterisation
330 can be performed. The Aerodyne Aerosol Mass Spectrometer (AMS) has the ability to
331 quantify the mass concentration and mass spectrum of organic matter giving insights into the
332 aerosol sources, chemistry and processes in the environment (Zhang et al. 2011).

333 Several studies have used an AMS for real-time measurement of particles and have identified
334 organic aerosols from cooking, referred to as Cooking Organic Aerosol (COA) (Mohr et al.,
335 2009; Allan et al., 2010; Huang et al., 2010; Mohr et al., 2011; Sun et al., 2011). Results from
336 this instrument provide real-time, highly time-resolved measurements of the concentrations
337 and size distributions of non-refractory submicron aerosol (NR-PM₁) species (i.e., organics,
338 sulphate, nitrate, ammonium, and chloride) with the capability to separate different organic
339 ions at the same nominal mass-to-charge ratio (m/z). The mass spectra (MS) represent the
340 linear superposition of the spectra of individual components weighted by their concentrations.
341 The mass information of the functional nature of the organics can be acquired on inspection
342 of the mass spectra obtained from the AMS after the removal of the inorganic contributions
343 (Allan et al., 2004; Allan et al., 2010). Multivariate factor analysis of the MS matrix has been

344 conducted in most studies and has provided OA factors that offer a quantitative generic
345 description of thousands of individual organic species (Zhang et al, 2011). The individual
346 factors typically correspond to a large group of OA constituents with similar chemical
347 composition and temporal behaviour that characterise different sources and atmospheric
348 processes. Nonetheless, the AMS provides limited molecular details of aerosols analysed
349 because the molecules are fragmented during high temperature vaporization and ionization in
350 the AMS (Canagaratna et al., 2007).

351 The main reference spectra of primary OA from cooking emissions (COA) derived from the
352 AMS have been obtained in Barcelona (Mohr et al. 2009, Mohr et al.2011), London (Allan et
353 al., 2010), New York and Beijing (Huang et al., 2010). A high resolution mass spectrum of
354 COA obtained in 2006 by Sun et al. (2011) was different from that reported by Mohr et al.
355 (2009), but similar to the one analysed in London and Beijing (Mohr et al., 2009; Allan et al.,
356 2010; Huang et al., 2010; Sun et al., 2011).

357 Characterisation of primary emissions from simulations of different types of Chinese cooking
358 and biomass burning was also done using a High Resolution Time of Flight AMS (HR-ToF-
359 AMS) at Peking University Shenzhen (He et al., 2010). The MS for all the cooking were
360 similar, with highest signals at m/z 41 and m/z 55 (dominated by $C_3H_5^+$ and $C_4H_7^+$), whilst
361 biomass burning MS showed the highest signals at m/z 29 and m/z 43. Therefore, there was a
362 significant difference between cooking and biomass burning mass spectra. On the other hand,
363 the MS of both cooking and biomass contained ions that demonstrated the presence of
364 saturated alkanes, alkenes, and long chain fatty acids in their primary OA (He et al., 2010).
365 The fragments that dominated Chinese cooking are believed to be fragments resulting from
366 electron ionization of unsaturated fatty acids from frying. The O/C ratios of OA ranged from
367 0.08- 0.13 for the Chinese cooking and 0.18 to 0.26 for biomass burning (He et al., 2010).

368 Another difference between cooking and biomass burning MS was the presence of more mass
369 fragments in the range of $m/z > 100$ for biomass burning than in the cooking emission profile.
370 He's study provides useful MS for the identification of individual components during factor
371 analysis of ambient OA datasets measured using an AMS.

372 Huffman et al., (2009) used a modified fast temperature –stepping thermodenuder coupled
373 with a HR-ToF-AMS to determine chemically resolved volatility of the OA emitted from
374 various primary sources of ambient OA, including meat cooking. Meat cooking OA was
375 found to have consistently lower volatility than ambient OA (Huffman et al., 2009). Meat
376 cooking MS was hydrocarbon-like and dominated by reduced ions (C_xH_y), with the abundant
377 ions being $C_3H_7^+$ (m/z 43) and $C_4H_7^+$ (m/z 55), whilst the contribution from oxygenated ions
378 was low. The range of O/C ratios was between 0.11-0.14 similar to those of typical fatty
379 acids like oleic acids, and much lower than for ambient urban Oxygenated OA (OOA)
380 (Huffman et al., 2009).

381 Prominent peaks were also identified in the study of Sun et al. (2011), which further
382 suggested that the suitable source spectral signature for cooking emissions and thus for use as
383 spectral markers for COA is the m/z ratio 55/57. The COA spectrum was characterized by a
384 high ratio of 2.9 and a high fraction of m/z 55 ($f_{55} = 8\%$), which confirmed the presence of
385 COA at urban locations (Sun et al., 2011). They also found that COA contributed 38% of
386 $C_3H_3O^+$ and 29% $C_4H_7^+$ at m/z 55; 25% of $C_3H_5O^+$ and 20% of $C_4H_9^+$ at m/z 57; and had a
387 significant correlation with a few $C_xH_yO^+$ ions (e.g., $C_5H_8O^+$, $C_6H_{10}O^+$, and $C_7H_{12}O^+$) (Sun et
388 al., 2011).

389 Mohr et al. (2009) reported that meat cooking aerosol had an elevated m/z 60 and contained
390 significant fractions of oxygenated organic compounds with similar mass spectral fractions
391 than ambient hydrocarbon like OA (HOA) (Mohr et al., 2009). Allan et al. (2010) identified

392 the largest peaks for COA at m/z 41 and m/z 55, similar to those identified by Lanz and
393 colleagues (2007) (Lanz et al., 2007). They also observed a diurnal profile which had large
394 peaks occurring in the late evening and smaller peak around at midday, which they ascribed
395 to mealtimes in London and Manchester (Allan et al., 2010).

396 Zhang et al. (2007) identified meat cooking as a part of the HOA after analysing organic
397 aerosol obtained in a field campaign using the AMS (Zhang et al., 2007). Combustion of
398 fossil fuel was the other source of HOA. Aerosol mass contributions from HOA in the urban
399 areas were strong, but smaller contributions were observed in rural areas attributed to
400 atmospheric dilution. OOA components had higher concentrations than advected HOA
401 indicating that oxidation of Secondary OA (SOA) was the strongest contributing source to
402 OOA, whilst HOA (and its various sources such as cooking) were less significant contributors
403 to organic aerosol (Zhang et al., 2007).

404 In California food cooking was found to contribute to 10% of submicron OA mass in the
405 summer periods, but due to high covariance, food cooking composition was not distinct from
406 the SOA during the autumn (Williams et al., 2010). During the summer the SOA had four
407 distinct components identified which accounted for about 88% of the total fine OA mass
408 when all were combined. The components included three distinct oxygenated sources: SOA1
409 (oxygenated species with hydrocarbons), SOA2 (oxygenated species from photo-oxidation of
410 gas-phase precursors), SOA3 (from advanced photochemical oxidation exist) and the fourth
411 component was from biogenic SOA.

412 Similarly, He et al. (2011) deployed a HR-ToF-AMS at an urban site in Hong Kong –
413 Shenzhen metropolitan area and the data obtained indicated four OA components which
414 included HOA, biomass burning (BBOA) and two oxygenated OA components with the

415 HOA contributing 29.5% of the total organic mass. Cooking emissions and fossil fuel
416 combustion were the identified sources of HOA (He et al., 2011).

417 AMS and ^{14}C data from Mexico City identified that non-fossil fuel sources (such as food
418 cooking and biogenic SOA) contributed to 38% of organic carbon and 28% total carbon
419 during the low regional fire period of their sampling (Aiken et al., 2010), further illustrating
420 the importance of cooking as a source contributing to aerosol concentrations in ambient air.

421 It is important to note the limitations of the AMS, which are that it only determines the non-
422 refractory material and does not distinguish between anthropogenic and biogenic secondary
423 organic aerosols (SOA) (Baltensperger and Prévôt, 2008). This can lead to both qualitative
424 and quantitative uncertainty in the reporting of data.

425

426 **4. SPECIFIC ORGANIC COMPOUNDS EMITTED DURING COOKING**

427 Cooking involves a wide range of chemical reactions. For instance, many sugars (e.g.
428 disaccharides or oligo-saccharides) or carbohydrates undergo hydrolysis when heated with
429 water. The hydrolysis reaction breaks down the complex sugar into single ring sugars. If
430 sugars are heated further, degradation reactions will occur and the sugar rings will open up to
431 form new molecules such as acids and aldehydes. If the temperature is increased sufficiently,
432 the degradation products may recombine to form chain-like molecules (Barham, 1950). In
433 meat cooking, fat which occurs as triglyceride (i.e. fatty acids esterified to a glycerol
434 backbone) in uncooked meat is hydrolysed or thermally oxidized and produces free glycerol,
435 free fatty acids and mono and diglycerides as shown in Figure 1 (Nolte et al., 1999). The
436 chemical reactions that occur between proteins and carbohydrates or sugars during cooking
437 are known as the Maillard reactions. These involve initial degradation to amino acids and

438 smaller sugars. The acids and aldehydes produced after the opening of the sugar rings react
439 with the amino acids to produce a wide range of chemicals (e.g. furanones) (Barham, 1950).

440 The chemical properties of the aerosols generated during cooking can be measured to further
441 provide useful information on the aerosol composition. In most of the studies aimed at
442 performing chemical speciation of the cooking aerosol, samples are collected on filters for
443 gravimetric determination and to allow subsequent chemical analysis. In some cases,
444 denuders are used to collect the vapour phase of semi-volatile components for further
445 analysis. Off-line chemical characterisation studies often employ sampling methods which
446 have the potential to cause positive artefacts associated with the reaction of trace gases with
447 particles on the filter or the filter itself. Negative artefacts may also arise from evaporative
448 loss of semi-volatile components. Strict sampling procedures and guidelines should keep
449 these artefacts to a minimum (Harrison and Yin, 2005).

450 Table 6 gives details of key studies that have sampled and subsequently analysed the
451 chemical composition of aerosols from cooking. Most of the off-line chemical
452 characterisation has been done using a GC-MS analytical stage for organic speciation of the
453 cooking emissions, with many compounds of interest requiring derivatisation. A summary of
454 specific groups of compounds emitted from cooking identified and characterised by these
455 studies appear in Table 7, whilst Table 8 to Table 18 present a summary of concentrations of
456 specific compounds emitted during cooking reported in the literature, as discussed below.

457

458 **4.1. Effect of cooking styles and ingredients on organic compound emission profiles**

459 Research has identified that different cooking styles emit different profiles of compounds.
460 The differences have been attributed to factors such as cooking processes and ingredients

461 (Hildemann et al., 1991a; Rogge et al., 1997; Schauer et al., 1999a; He et al., 2004a).
462 Western fast food cooking involves frying with beef and chicken as the main cooking method
463 and meats consumed. Chinese cooking practice on the other hand generally involves the use
464 of pork, poultry, seafood as well as vegetables during cooking as listed in Table 1. Chemical
465 composition variations are thus expected to be observed for various different cooking
466 operations. For instance, nonanedioic acid has been identified as the most abundant
467 dicarboxylic acid in Chinese cooking and hexanedioic acid for meat cooking (Rogge et al.,
468 1991; He et al., 2004c; Zhao et al., 2007c). Sitosterol and monosaccharide anhydrides have
469 been attributed to vegetables used in Chinese cooking as they were not observed in meat
470 cooking processes. These differences in chemical composition need to be considered for
471 selection of molecular markers, which will be useful to assess the contribution of cooking to
472 atmospheric particulate organic matter (POM) (Rogge et al., 1991; He et al., 2004b; Zhao et
473 al., 2007c; Zhao et al., 2007b). Figure 2 shows Marker-to-OC ratios of meat cooking profiles
474 using profiles from Rogge et al. (1991), Watson et al. (1998) and Schauer et al. (1999a;
475 Schauer et al., 2002). These source profiles and species are usually included in models by
476 normalising emissions with OC or PM_{2.5}.

477 Higher fat contents in cooking ingredients have been found to produce more fatty acids
478 compared with the low fat content ingredients in the same cooking operation (Rogge et al.,
479 1991; Zhao et al., 2007b; Zhao et al., 2007c). This is generally observed also when Chinese
480 cooking is compared with Western style fast food; the latter having higher concentrations of
481 fatty acids, indicating the high proportion of ingredients with higher fat content. Animal and
482 vegetable fats are rich in high concentrations of normal fatty acids with even carbon numbers
483 from C4 to C34 as triglycerides and phospholipids (Zhao et al., 2007a).

484 In a experiment comparing grilling and charbroiling different types of meat, grilling was
485 found to emit less organic compounds than charbroiling, which yielded about 5 times more
486 PAH (i.e. 30-50 mg/kg for charbroiling vs. <10 mg/kg for grilling), 10 times more lactones
487 (i.e. 7-30 mg/kg for charbroiling vs. 2-4 mg/kg for grilling) and 20 times more cholesterol
488 (i.e. 1-8 mg/kg for charbroiling vs. 0.04-0.2 mg/kg for grilling) (McDonald et al., 2003) .

489 When different types of meat were grilled in a shed, Mohr et al. (2009) reported large
490 differences of emissions with increasing emissions as the fat content increased, even when
491 the meats were cooked in the same manner. This is qualitatively consistent with earlier
492 studies (Rogge et al., 1991; McDonald et al., 2003). Rogge et al. (1991) reported that
493 generally grilling of meat led to the higher production of aerosols made of fatty acids. This
494 was attributed to the oil and grease droplets falling into the gas flame or onto the heat source
495 where they would vaporize and renucleate and grow into small particles.

496 Zhao et al. (2007a) investigated the chemical composition of particulate organic matter from
497 Western fast food cooking and identified tetradecanoic acid, hexadecanoic acid, octadecanoic
498 acid, 9-octadecanoic acid, nonanal, levoglucosan, hexanedioic acid and nonanedioic acid as
499 potential tracers with saturated and unsaturated fatty acids accounting for 78% of total
500 quantified compounds. When they analysed the chemical composition of aerosol from
501 Chinese cooking, they identified also a dominant homologue of fatty acids with its
502 concentration being about 73-85% of the quantified compounds. They also identified
503 levoglucosan and β -sitosterol as well as a clear pattern of n-alkanes which were taken as an
504 indication of vegetables consumed in the Chinese cooking process (Zhao, 2007b). The
505 concentration of quantified compounds per unit of particulate organic matter in Western
506 cooking was found to be much higher than that in Chinese cooking (Zhao, 2007a). The
507 candidate organic tracers that they found to distinguish emissions of Western cooking from

508 Chinese cooking in Ghanzou (China) are tetradecanoic acid, hexadecanoic acid, octadecanoic
509 acid, oleic acid, levoglucosan, mannosan, galactosan, nonanal and lactones (Zhao, 2007b).
510 Table 8 shows clearly from their findings that the Chinese cooking made a much greater
511 contribution of PAHs to particulate organic matter than Western fast food with 2,855 ng/mg
512 of particulate organic matter in Chinese cooking as against 40 ng/mg in Western cooking.

513 Nolte et al. (1999) analysed meat cooking smoke and found that 1-palmitin and 2-palmitin
514 were the most abundant compounds observed with significant emissions of 1-stearin and 1-
515 olein monoglycerides and cholesterol (Nolte et al., 1999). Similar to what was observed with
516 emissions of particulate number and particulate matter mass, higher concentrations of organic
517 pollutants were observed to be emitted during oil-based cooking methods compared to
518 steaming and boiling which were water-based (See and Balasubramanian, 2008). Also an
519 analysis of commonly used cooking fuels in Hong Kong identified that gas cooking produced
520 higher concentrations of PM₁₀, organic material and total volatile organic compounds during
521 cooking by stir frying, pan frying and deep frying in domestic settings (To and Yeung, 2011).
522 Higher concentrations of pollutants were observed in commercial kitchens compared to
523 domestic kitchens probably due to the volume of food cooked or methods of cooking used. In
524 the commercial restaurant, broiling of meat was found to produce higher concentrations of
525 PM and VOC especially for electric broiling of meat compared to gas broiling. This was
526 attributed to a larger contact area for the beef on the electric broiler compared to the gas
527 broiler leading to more intense effect of the heat (To and Yeung, 2011).

528

529 **4.2. Emissions of Polycyclic Aromatic Hydrocarbons**

530 The incomplete combustion or pyrolysis of organic substances containing hydrogen and
531 carbon, leads to the production of polycyclic aromatic hydrocarbons (PAH). They are stable

532 organic molecules, some of which are known carcinogens (USEPA, 1999; IARC, 2010).
533 Table 9 summarises the concentrations of PAH in fumes reported in the literature with
534 different cooking styles, methods and ingredients; whilst Table 10 summarises the
535 concentrations in fumes from different oils. The cooking method used has been identified as
536 one of the influential factors affecting emission of some compounds, with deep frying found
537 to produce more pollutants and an abundance of higher molecular weight PAHs.

538 Li et al. (2003) analysed fumes from the rooftop exhausts of ten restaurants producing
539 Chinese, Western, fast food, and Japanese types of food in Taiwan and studied their PAH
540 content. They found that Chinese cooking contributed higher levels of benzo[a]pyrene
541 equivalents than all other styles of cooking, with Western food contributing about seven
542 times less than the Chinese method. Japanese and Western fast food showed a negligible
543 contribution to benzo[a]pyrene. This is consistent with later work by Zhao et al. (2007c), who
544 did not detect any PAH in Western style fast food cooking.

545 Nonetheless, the percentage of PAH in Chinese cooking as a proportion of total quantified
546 compounds is generally small in emissions from cooking. The compounds of highest
547 concentration emitted from Chinese cooking are pyrene (He et al., 2004b) with traces of
548 benzo[a]pyrene present (Zhao et al., 2007b). Meat and seed oil cooking were observed to
549 release mainly chrysene (Rogge et al., 1991; Schauer et al., 1999a; Schauer et al., 2002).
550 Samples collected in the breathing area of chefs working at Norwegian à la carte restaurants
551 contained naphthalene within the range of 0.05-0.27 $\mu\text{g}/\text{m}^3$ with a total mean value for the
552 restaurants being 0.18 $\mu\text{g}/\text{m}^3$ (Sjaastad and Svendsen, 2009).

553 The relatively low proportion PAH compounds in Chinese emissions was also observed by
554 See et al. (2006) when they made a health risk assessment of occupational exposure to PAH
555 associated with Chinese, Malay and Indian cooking. They found that Malay cooking emitted

556 higher PAH concentrations and also a larger fraction of PAH in PM_{2.5} (600 ng/m³ and 0.25%,
557 respectively) than the other methods (Chinese, 141 ng/m³ and 0.07%; Indian, 37.9 ng/m³ and
558 0.02%, respectively). This was attributed to extensive frying of snacks (See et al., 2006).
559 Deep frying generated more PAH than other cooking methods due to the high temperature
560 during cooking as well as the large amount of oil used for this method of cooking.

561 This trend was further observed when Chinese cooking and Indian cooking were compared:
562 higher PAH concentrations were observed for Chinese cooking due to stir frying and higher
563 cooking temperature, whilst the Indian cooking style generated the lower PAH
564 concentrations. Indian cooking emitted large amounts of volatile PAH with lower molecular
565 weight like naphthalene, fluoranthene and phenanthrene attributed to low temperature
566 cooking, such as simmering (See et al., 2006). Chinese cooking, on the other hand, was found
567 to emit higher molecular weight PAHs such as benzo[b]fluoranthene, indeno[1,2,3-cd]pyrene
568 and benzo[g,h,i]perylene. These trends were attributed to the cooking methods employed in
569 each type of cooking from the amount of food cooked, the amount and type of oil used, to the
570 temperatures reached during cooking, and cooking time (See et al., 2006).

571 The effect of the cooking method was also examined by See and Balasubramanian (2008),
572 who found that techniques that involve the use of oil at high temperatures, such as stir frying,
573 pan-frying and deep-frying, released higher amount of PAH compared with those that involve
574 the use of water, such as boiling and steaming. This is consistent with work of Schauer et al.
575 (2002). Higher quantities of oil are generally used in stir frying, commonly used in Malay
576 and Chinese cooking, than simmering which is the most common technique used for
577 preparation of Indian dishes (Table 1). In addition, high temperature frying was found to lead
578 to production of higher molecular weight PAHs, while low temperature cooking results in
579 formation of more low molecular weight PAHs (See et al., 2006). McDonald et al. (2003)

580 compared the PAH emissions from charbroiling and grilling meat and found that PAH
581 emissions from charbroiling were about 3–5 times more than when food was grilled. This
582 was attributed to the contact of the lipid material dripping from the meat (during cooking)
583 onto the cooking appliance. Thus, the higher PAH concentrations observed during
584 charbroiling were due to the direct access of lipids onto the hot flame compared to the cooler
585 griddle surface used in grilling (McDonald et al., 2003).

586 The emission of PAHs in cooking fumes, not only is related to the cooking method, but also
587 to the cooking ingredients. Schauer et al. (1999a; 2002) studied the emissions of cooking
588 fumes for charbroiling hamburger meat (1999a) and frying vegetables (2002). They found
589 that cooking meat produced far greater PAH concentrations than frying vegetables. Zhu and
590 Wang (2003) studied the emissions of low and high fat food using different cooking methods.
591 The frying of low fat foods was observed to lead to the generation of more PAH than the
592 broiling. This was not the case for high fat food which exhibited the reverse with higher
593 concentration of PAH detected when the food was broiled (Zhu and Wang, 2003). This was
594 illustrated when low fat fish produced a higher level of PAH when fried than when broiled,
595 and pork chops produced higher PAH when broiled than when fried.

596 In another study of cooking in domestic and commercial kitchens it was observed that PAH
597 concentrations were higher for increased cooking temperatures with mean measured
598 benzo[a]pyrene concentrations of 6-24 ng/m³ in domestic kitchens and 150-440 ng/m³ in
599 commercial kitchens (Zhu and Wang, 2003). The difference was attributed to the large
600 amount of food cooked in the commercial kitchen.

601 The type of oil used for cooking will also affect the composition of cooking fumes and its
602 PAH content as summarised in Table 10. Fortmann et al. (2001) analysed five untreated (i.e.
603 raw and uncooked) cooking oils (canola, olive, corn, peanut and vegetable oils) for PAH in a

604 controlled environment. Olive oil and peanut oil were found to contain higher concentrations
605 of PAH than the other oils analysed (Fortmann et al., 2001). Chiang et al. (1999) carried out
606 analysis on the fumes emitted from lard oil, peanut oil and soya bean oil by and found that
607 fumes from these commonly used commercial cooking oils in Taiwan contained
608 dibenzo[a,h]anthracene and benzo[a]anthracene. Benzo[a]pyrene was also found in fumes in
609 peanut oil and soya bean oil. Chiang et al. (1999) found that extracts of fumes from
610 sunflower, vegetable and corn oils contained all the above PAHs as well as
611 benzo[b]fluoranthene with sunflower oil fumes having high concentrations of benzo[a]pyrene
612 of about $22.7 \mu\text{g}/\text{m}^3$ and corn oil the least, $18.7 \mu\text{g}/\text{m}^3$ (Chiang et al., 1999).

613 Wu et al. (1998) analysed the fumes from three different commercial cooking oils (lard oil,
614 soya bean oil and peanut oil) frequently used in Chinese cooking and found that fumes
615 contained PAHs, nitro-PAHs such as 1-nitropyrene and 1,3-dinitropyrene (Wu et al., 1998).
616 In another study it was observed that cooking with lard emitted higher concentrations of
617 PAHs, followed by soya bean oil and rapeseed oil (Zhu and Wang, 2003). Chen and Chen
618 (2001) analysed the polycyclic aromatic hydrocarbon (PAH) contents of smoke emitted from
619 selected cooking oils and they observed that soybean oil generated a larger amount of PAH
620 than canola oil or sunflower oil. In a subsequent study where they examined the smoke
621 emitted during the cooking of chicken legs they found that canola oil emitted the greatest
622 content of PAHs, followed by soybean and sunflower oil (Chen and Chen, 2001; Chen and
623 Chen, 2003). This contradictory observation of soybean oil or canola oil releasing more PAH
624 contents in the two studies was attributed to the heating of oil with or without food
625 ingredients as well as possible difference in smoke collection device used in the different
626 experiments. Chen and Chen (2003) also found that a much greater content of smoke was
627 generated when the oils were heated with chicken leg than when heated with glass beads
628 alone.

629 4.3. Emissions of n-Alkanes

630 Normal alkanes have been found to range from C₁₄ to C₃₃ for the various cooking techniques
631 (He et al., 2004b; Zhao et al., 2007c) as summarised in Table 11. The distribution of n-
632 alkanes emitted from Chinese restaurants have generally been observed to be substantially
633 different from the distribution from meat cooking (Rogge et al., 1991; Schauer et al., 1999a;
634 He et al., 2004b) and similar to alkane patterns from frying vegetables in seed oils (Schauer et
635 al., 1999a; Schauer et al., 2002). Emission of n-alkanes from cooking consisted of a
636 negligible fraction of the total quantified organic mass emitted and is dependent on the
637 cooking conditions (Rogge et al., 1991; He et al., 2004b). Hildemann et al., (1991a) reported
638 that the n-alkane concentration release rate increased from frying to charbroiling of meat with
639 extra lean meat releasing less compounds than regular meat (Hildemann et al., 1991a). This
640 was similar to observations by Rogge et al., (1991), where charbroiling was found to produce
641 three times the mass of n-alkanes than frying of meat (16 mg/kg of charbroiling meat as
642 against 5.5 mg/kg of frying meat). Rogge et al. (1991) also observed that charbroiling regular
643 meat released four times the mass compared to extra lean meat (thus affected by fat content
644 of meat).

645 Western style fast food cooking has been observed by Zhao and co-workers to emit double
646 the concentration of n-alkanes per mg particulate organic matter (POM) compared to Chinese
647 cooking. They also observed that Chinese cooking has an odd to even carbon preference
648 when carbon number is greater than 23, whilst Western style fast food does not (Zhao et al.,
649 2007a). This is taken as an indication of the high plant wax content of Chinese cooking (Zhao
650 et al., 2007b). The n-alkanes have a C_{max} at C₂₅ for western fast food (Zhao et al., 2007a) and
651 meat cooking (Rogge et al., 1991). Chinese cooking exhibits a C_{max} at C₂₉ or C₃₁ taken as an
652 indication of the presence of vegetables during cooking operations.

653 4.4. Emissions of Fatty Acids

654 Meat and oils used in cooking contain fats made up of saturated and unsaturated fatty acid
655 esters of glycerol. The chemical processes that typically occur during high temperature
656 treatment of food are the degradation of sugars, pyrolysis of proteins and amino acids and the
657 degradation of fats (Svendsen et al., 2002). The cooking process leads to production of free
658 fatty acids, free glycerol and mono- and diglycerides (Nolte et al., 1999). Table 12
659 summarises the concentrations of fatty acids reported in the literature.

660 In Western fast food cooking, the quantified saturated fatty acids observed range from C₆ to
661 C₂₀ with distinct even to odd carbon preference and a predominance of palmitic acid (Zhao et
662 al., 2007a). Chinese cooking was found to emit C₆-C₂₄ fatty acids with a similar even to odd
663 carbon preference and palmitic acid preference similar to meat cooking (Rogge et al., 1991;
664 He et al., 2004b) and seed oil cooking (Schauer et al., 2002). The most common unsaturated
665 fatty acids observed were oleic acid and linoleic acid for Chinese cooking (Zhao et al.,
666 2007b; He et al., 2004b). The most prominent organic compound released from American
667 cooking is oleic acid (Rogge et al., 1991; Schauer et al., 1999a; Schauer et al., 2002; He et
668 al., 2004b).

669 The concentration of emitted saturated fatty acids in western fast food was found to be 13
670 times higher than in Chinese cooking while unsaturated fatty acid concentrations were only
671 two times higher, attributed to ingredients and cooking temperature. The ratio of oleic acid to
672 stearic acid in western style fast food cooking is less than one, but greater than one in Chinese
673 cooking. This was attributed to the high temperature in fast food which favours degradation
674 of triglyceride and phospholipids to release and evaporate fatty acids and reduce the
675 production of unsaturated acids (Rogge et al., 1991; Zhao et al., 2007a). High concentrations
676 of nonanoic acid emissions are observed in both Chinese and Western style fast food cooking

677 with a higher ratio of nonanoic acid to other acids (C₈-C₁₀) in Western style fast food.
678 Schauer et al. (1999a; Schauer et al., 2002) compared the emissions of fatty acids from
679 different ingredients, such as meat and vegetables. They found that charbroiling hamburger
680 meat released more fatty acids than frying vegetables. They also found that stir frying
681 released more fatty acids than deep frying.

682 Peanut oil, commonly used in Chinese cooking, was heated in a wok and heated to a
683 temperature of 260°C. Gas phase organic emissions were found to contain several n-alkanoic
684 acids as methyl esters of hexadecanoic acid (palmitic acid), oleic acid (octadecenoic acid)
685 acid and dicarboxylic acid- linoleic acid (octadecadienoic acid). It was concluded that the
686 fatty acids were derived from the tri-acylglycerides and di-acylglycerides of peanut oil (To et
687 al., 2000). These further highlighted the importance of temperature in cooking as the
688 temperature used in the study (260°C) was not sufficient to lead to the complete breakdown
689 of the fatty acid components (To et al., 2000).

690

691 **4.5. Emissions of Dicarboxylic Acids**

692 Dicarboxylic acids are released from the oxidation of dialdehydes during autoxidation of
693 unsaturated lipids and are generally measured in the range of C₄-C₁₁ (Rogge et al., 1991; He
694 et al., 2004b). Hexanedioic acid followed by nonanedioic acid, have been identified as the
695 predominant acids in Western fast food cooking (Zhao et al., 2007a). Nonanedioic acid is the
696 predominant dicarboxylic acid identified from Chinese cooking (He et al., 2004b; Zhao et al.,
697 2007b). Meat cooking showed highest concentration for hexanedioic acid (Rogge et al., 1991,
698 Schauer et al., 1999a) and the study of cooking with seed oil showed a predominance of
699 hexanedioic acid and octanedioic acids (Schauer et al., 2002). A higher concentration of this
700 acid has been observed in western fast food with the amount of dicarboxylic acid per mg of

701 POM in Chinese cooking being about nearly 30 times less than in Western cooking (Zhao et
702 al., 2007a). This has been attributed to the breakdown of unsaturated fatty acids and
703 unsaturated lipids (Zhao et al., 2007a). A summary of the concentrations of dicarboxylic
704 acids reported in the literature is found in Table 13.

705

706 **4.6. Emissions of Aldehydes**

707 A recent IARC monograph reported that cooking, in particular frying, generates substantial
708 amounts of certain gaseous pollutants such as formaldehyde (IARC, 2006), acetaldehyde
709 (IARC, 1999), acrylamide (IARC, 1994) and acrolein (IARC, 1995).

710 Concentration distributions have been found to be similar for Western and Chinese style
711 cooking for most of the aldehydes (Table 14), except for nonanal, which is one order of
712 magnitude higher in Western cooking (Zhao et al., 2007b).

713 Similar to what was observed for other organic species, the type of ingredient cooked is also
714 key in the release of aldehyde emissions during cooking. The studies of Schauer et al. (1999a;
715 Schauer et al., 2002) show that charbroiling hamburger meat emits more aldehydes than
716 frying vegetables.

717 Sjaastad et al. (2010) used a model kitchen similar to a Western European restaurant to assess
718 if higher mutagenic aldehydes were emitted during the frying of beefsteak on an electric
719 or gas stove with margarine or soya bean oil as the frying fat oil. It was found that mutagenic
720 aldehydes were detected in the breathing zone of cooks in the range of non-detectable to
721 $61.80 \mu\text{g}/\text{m}^3$ (Sjaastad et al., 2010). They also found that higher exposures to these
722 components were more pronounced when frying on a gas stove instead of an electric stove
723 which may cause adverse health effects especially for people occupationally exposed to these

724 fumes (Sjaastad et al., 2010). An earlier study of Sjaastad and Svendsen (2008) had involved
725 the frying of beef steak using margarine, rapeseed oil, soybean oil or virgin olive oil as frying
726 fat in similar conditions as a regular Norwegian home (in terms of ventilation conditions and
727 frying procedure). They recorded mutagenic aldehyde concentrations ranging from non-
728 detectable to 25.33 $\mu\text{g}/\text{m}^3$ (Sjaastad and Svendsen, 2008). They also observed statistically
729 significantly higher levels of mutagenic aldehydes and particles when margarine was used as
730 the cooking fat compared to the other oil.

731 Amides are formed during cooking (Stadler et al., 2002). N-glycosides formed by reaction of
732 reducing sugars with asparagines - one of the common natural amino acids- when heated,
733 resulted in significant levels of acrylamide (Figuerola et al., 2006). Rogge et al. (1991)
734 reported the presence of several amides in smoke released during charbroiling hamburger
735 meat, such as palmitamide, stearamide, and oleamide. Another nitrogen-containing
736 compound found was N,N-dibutylformamide. Frying hamburgers emits 2.8 mg of amides/kg
737 of meat, whereas gas broiling extra-lean meat increases the amount released by 13-fold (36
738 mg/kg of meat); doubling the fat content (regular meat) further doubles the amount emitted
739 (70 mg/kg of meat) (Rogge et al., 1991). Zhao et al. (2007c) compared the emissions of
740 several amides in the smoke released during Western and Chinese cooking, and found that
741 Western style fast food cooking released larger quantities of amide compounds.

742

743 **4.7. Emission of N-alkanones and Lactones and Organic Ions**

744 N-Alkanones (Table 15) are also detected from cooking, with 2-pentadecanone the most
745 prominent alkanone. Concentration distributions have been found to be similar for western
746 and Chinese style cooking (Zhao et al., 2007b) for most ketones, although 2-pentadecanone,

747 2-hexadecanone and 2-heptadecanone are orders of magnitude higher in Western style
748 cooking.

749 Lactones (Table 15) are detected from emissions of Western style fast food, meat
750 charbroiling, seed oil and Chinese cooking operations (Zhao et al., 2007a; Zhao et al., 2007b;
751 Schauer et al., 2002; Rogge et al., 1991). They are not emitted in large quantities, but can
752 serve as good cooking tracers. The Western cooking style releases more lactones than
753 Chinese cooking. For both n-alkanones and lactones, and similar to previous trends observed
754 for other organics, temperature during cooking has also been shown to have an influence on
755 their release, with emissions from stir frying higher than those from deep frying (Schauer et
756 al., 2002).

757 See and Balasubramanian (2008) reported the content of organic ions in the smoke produced
758 when using several cooking techniques. Consistent to the trend observed for other organics,
759 techniques that involve the use of water, such as steaming and boiling, released lower amount
760 of organic ions than those which involve the use of hot oils. Nonetheless, for organic ions,
761 deep-frying releases the highest amount of organic ions (Table 16).

762

763 **4.8. Emissions of Molecular Markers**

764 These are organic compounds occurring naturally in food which can be used as tracers. Some
765 major biomarkers for food are cholesterol, sterols and monosaccharide anhydrides, as
766 summarised in Table 17. Sterols are widely present in animal and vegetable body tissues
767 with plant lipid waxes and membranes generally comprised of C₂₈ and C₂₉ pythosterol
768 compounds such as β -sitosterol. Cholesterol is biosynthesised by higher animals and found in
769 body tissues such as animal fats and oils (Rogge et al., 1991; Zhao et al., 2007b).

770 Molecular markers identified for Chinese and Western cooking are levoglucosan, galactosan
771 and cholesterol (Zhao et al., 2007a; Zhao et al., 2007b; He et al., 2004b) while only
772 cholesterol has been detected for meat cooking (Schauer et al., 1999a; Rogge et al., 1991).
773 Schauer et al. (1996), Schauer and Cass (2000) and Schauer et al. (2002) have also identified
774 cholesterol as a tracer for activities of meat cooking.

775 Combinations of molecular biomarker compounds may be useful in distinguishing cooking
776 source type. Zhao et al. (2007a) combined concentrations of cholesterol and levoglucosan
777 with other compounds such as β -sitosterol, hexanoic acid and nonanedioic acid and were
778 able to differentiate emissions from western style fast food from Chinese cooking. This was
779 done by considering the ratio of meat to vegetable in both cooking styles as well as the
780 differences in concentration ratios of different chemical species.

781 The percentages of levoglucosan and cholesterol to total quantified compounds in Western
782 style fast food were observed to be lower than in Chinese cooking indicating that cooking oil
783 may have an effect on the total emission of compounds (Zhao et al., 2007a).

784

785 **4.9. Emissions of Elemental Carbon and Inorganic Compounds**

786 Cooking fumes contain carbonaceous particles (containing elemental carbon (EC) as well as
787 organic carbon (OC)) mainly within the fine particle fraction of the emission (See and
788 Balasubramanian, 2008). Organic carbon is the major constituent found in cooking fumes
789 (Kleeman et al., 1999; Schauer et al., 1999a; Chow et al., 2004; Kleeman et al., 2008; See
790 and Balasubramanian, 2008). Comparison of different culinary methods (Table 18) showed
791 that those using oil released higher concentrations of EC and OC than those using boiling (i.e.
792 boiling and steaming). EC and OC follow a similar trend to that observed with the release of

793 organic ions, with the highest emissions attributed to deep frying (See and Balasubramanian,
794 2008).

795 Particle bound water soluble ions have also been identified in cooking fume exhaust Na^+ , K^+ ,
796 NO_2^- , Mg^{2+} , Ca^{2+} , Cl^- , NO_3^- , and SO_4^{2-} (Hsieh et al., 2011). See and Balasubramanian (2008)
797 found that no major differences were observed between different cooking techniques (Table
798 18).

799 The metal content of the aerosol emitted during cooking was analysed by See and
800 Balasubramanian (2008) during steaming, boiling, stir-frying, pan-frying and deep frying.
801 They found that generally, the higher concentrations were observed in those techniques that
802 used mainly oil, and frequently the highest concentrations were found in the smoke released
803 during deep-frying (Table 18).

804

805 **5. APPORTIONMENT STUDIES**

806 Source apportionment studies are conducted to identify sources of pollution and the
807 contributions from the various sources to total pollution; eliminating the tendency to
808 emphasize certain sources while ignoring others. Source apportionment aids in the
809 identification of major sources which can be useful in the development of best possible
810 control practices to ensure adequate reduction of pollution.

811 Early source apportionment studies used mainly data for elements such as Fe, Zn, Pb, Cr, Al
812 and Ni together with inorganic ions as tracers for source identification and as there are
813 usually various sources of the same element it often has proved difficult to identify pollution
814 sources with confidence (Pant and Harrison, 2012). This problem has been reduced with the
815 use of organic compound markers (Lin et al., 2010). Organic compounds are generally

816 representative of their particular pollution sources and their selection for use in receptor
817 models depends on a) the chemical species selected to characterise the source should not be
818 reactive and should be conserved in the atmosphere; b) the source profiles should be linearly
819 independent and c) the uncertainties of measurement should be random, uncorrelated, and
820 normally distributed (Lin et al., 2010).

821 Receptor modelling of airborne particulate matter involves the application of multi-
822 component chemical composition data to attribute the mass measured in the atmosphere to
823 specific sources of emission. Ambient pollutant measurements are collected at sampling sites
824 to deduce the information on the sources of pollution and how it affects concentrations in the
825 location. Measurements of well as chemical species generated during cooking processes
826 provide a profile for the cooking emissions which in combination with the profile of other
827 sources such as traffic are a useful tool in source profile identification.

828 The most commonly used receptor model to apportion cooking sources is the Chemical Mass
829 Balance (CMB) model. Other source apportionment studies on cooking emissions have used
830 statistical techniques such as the Diagnostic Ratio, Principal Component Analysis (PCA) and
831 Positive Matrix Factorisation (PMF) for understanding the sources of PM emitted to the
832 atmosphere (Hopke et al., 2003; Robinson et al., 2006; See and Balasubramanian, 2006b;
833 Zhao et al., 2006; Kleeman et al., 2008; Brinkman et al., 2009; Allan et al., 2010; Clougherty
834 et al., 2011).

835

836 **5.1. Chemical Mass Balance (CMB) Model**

837 This model requires chemical source profiles which describe the specific chemical
838 composition of emitted particles as input in order to quantify those sources within the data

839 and apportion for the sources in the atmosphere. Source profile and fitting specie selection for
840 CMB analysis is a sensitive process which requires careful consideration as the profiles must
841 be adequately different for all the sources included in the model to ensure proper
842 apportionment at the receptor. The source emissions in the profile should not interact with
843 each other during transport and also their chemical and physical properties should be
844 practically constant during their transport from source to receptor (Chow and Watson, 2002).
845 Some sources do not have existing source signatures or ones specific to the location being
846 studied. In such cases profiles are borrowed from other cities with similar pollution sources,
847 which may not represent the source of emissions in the sampling area of interest perfectly
848 (Lee et al., 2008b; Pant and Harrison, 2012). Similarly when source profiles are too similar,
849 the CMB model yields large uncertainties in source contributions (Chow and Watson, 2002).
850 Generally the species used in most CMB models are from the source profiles available
851 through the USEPA Speciate database. The source profiles consist of both organic and
852 inorganic aerosol constituents (Schauer and Cass, 2000). The source profiles generally used
853 for charbroiled meat cooking, gasoline vehicle emissions, diesel truck emissions and paved
854 road dust (Hildemann et al., 1991a; Schauer et al., 1999a; Schauer et al., 1999b; Fraser et al.,
855 2002) and for vegetative detritus and natural gas combustion (Rogge et al., 1993a; Rogge et
856 al., 1993b) are mainly obtained from studies in the United states such as Texas and Los
857 Angeles (Zheng et al., 2002; Fraser et al., 2003). However, the use of source profiles from
858 locations other than the area of study might introduce uncertainty in the apportionment of
859 some source contributions (e.g. road dust; soil).

860 A review by Lin et al., 2010 identified the need for a more specified organic compound
861 markers for some particle sources such as non-meat cooking particle emissions, paved roads,
862 fugitive dust, biogenic, and agriculture emissions as well as a source contributions library for
863 particular locations, for use in CMB models.

864 Mass balance models that have been applied in indoor environments have usually taken into
865 consideration various combustion related activities like home heating and cooking; and also
866 activities such as cleaning and infiltration of outdoor air resulting in a contribution from
867 outdoor sources (Millar et al., 2010). CMB analyses have made use of different combinations
868 of source profiles for the estimation of the contribution of food cooking emissions to ambient
869 particle concentrations. Several food cooking source profiles have been published (Rogge et
870 al., 1991; Nolte et al., 1999; Schauer et al., 1999a; Rogge, 2000; Schauer and Cass, 2000;
871 Chow et al., 2004; Robinson et al., 2006). These cooking profiles contain speciated organic
872 data with a range of emission composition and rates mainly dependent on cooking technique
873 and food type. The use of source profiles and fitting species require that the model must
874 include all major sources and the species should be conserved during transport from source to
875 receptor (Watson et al., 1998; Robinson et al., 2006). Organic molecular markers such as
876 oleic acid, cholesterol and palmitic acids are used to estimate the contribution of food
877 cooking emission to primary organic aerosol (Rogge et al., 1991; Schauer et al., 1999a;
878 Robinson et al., 2006). Cooking has in some cases been classified with agricultural burning
879 and wood smoke due to emissions of similar organic compounds from the processes (e.g.
880 levoglucosan and cholesterol) (Chow and Watson, 2002).

881 Using the chemical mass balance model to apportion for the sources of $PM_{2.5}$ in a city has
882 been exemplified by Schauer et al. (1996). They found that the organic carbon mass
883 contribution of PM due to meat cooking was about 23% in Los Angeles, which was
884 comparable to findings by Hildemann et al. (1991b) and Rogge et al. (1991) in earlier studies.
885 The CMB approach was also used to find that meat cooking contributed between 20% and
886 75% to ambient concentrations of PAHs in residential areas (Venkataraman and Friedlander,
887 1994).

888 Robinson et al. (2006) made use of the basic set of source classes and compounds developed
889 by Schauer et al. (1996) and Schauer and Cass (2000). The CMB analysis included source
890 profiles of eight source classes: diesel vehicles, gasoline vehicles, road dust, biomass
891 combustion, cooking emissions, coke production, vegetative detritus and cigarettes. However,
892 Robinson et al. (2006) reported that a large systematic bias was generally observed in CMB
893 models due to differences in species and source profile marker to organic carbon ratios. The
894 ambient ratio of palmitic acid to oleic acid was higher than expected from other published
895 literature, reflecting problems presented by source profile variability. This signified that the
896 CMB could not fit both the acids simultaneously, even though ambient concentrations
897 showed a strong correlation indicating they were from the same source. The use of the two
898 alkanolic acids as fitting species in the model in addition to other cooking markers however
899 provided a better model for source contribution estimates, further highlighting the importance
900 of molecular markers in source apportionment analysis. The model apportioned 320 ± 140 ng
901 C/m^3 (10% of the study average ambient organic carbon) to food cooking emissions.

902 In the south-eastern United States, particle phase organic compounds were used in a CMB
903 model to understand how the primary source contributions of particulate matter and fine
904 particulate organic carbon concentrations varied with seasons (Zheng et al., 2002). The
905 results indicated that wood smoke, meat cooking and gasoline powered motor vehicles
906 contributed to $PM_{2.5}$ organic carbon concentrations in the range of 25-66%, 5-12% and 0-
907 10% respectively, with minor contributions from paved road dust and vegetative debris.
908 Between 2003 and 2004, Zheng et al. (2002) sampled again four sites of the Carbonaceous
909 Aerosol Characterization Experiment (CACHE) and used CMB and carbon isotope analysis
910 to further understand variability of organic components and source contributions to fine
911 organic carbon and $PM_{2.5}$ in the south-eastern United States. Meat cooking was again
912 identified as a primary emission source of OC along with eight other sources including wood

913 combustion (which was the most dominant source, 14-23%), gasoline engine exhaust, diesel
914 engine exhaust, vegetative debris, cigarette smoke, road dust and natural gas exhaust (Zheng
915 et al., 2006). The carbon isotope analysis showed results consistent with the CMB analysis,
916 with both identifying that urban areas had a high fossil fraction of carbon.

917 Meat cooking operations were also identified as one of the sources of ambient fine particulate
918 matter in Houston Texas with a contribution of between 0.9-1.3 $\mu\text{g}/\text{m}^3$ at urban sites and 0.7
919 $\mu\text{g}/\text{m}^3$ at background sites (Fraser et al., 2003). Other important sources identified were diesel
920 exhaust (1.6- 3.7 $\mu\text{g}/\text{m}^3$ at the urban site), gasoline powered vehicles, paved road dusts and
921 wood combustion. These sources were identified using eight source profiles in the model
922 which used 24 molecular markers.

923 CMB analysis of organic molecular marker data in Pittsburgh Pennsylvania also identified
924 cooking as an anthropogenic source of organic aerosol and $\text{PM}_{2.5}$ and found that secondary
925 organic aerosols were actually the major components of organic carbon (OC) in Pittsburgh in
926 all seasons, whilst primary sources affected ambient concentrations only occasionally
927 (Subramanian et al., 2007). This is unlike other locations where the contributions of meat
928 cooking and traffic are unaffected by season and show no seasonal variation (Fraser et al.,
929 2003).

930 In Atlanta meat cooking was among the major contributors of fine OC identified with a range
931 of 7-68% (average 36%) in summer periods and 1-14% (average 5%) during the winter
932 months. Gasoline and diesel exhaust contributed 21% and 20% respectively to OC during the
933 summer and 33% to 4% during the winter, with wood combustion being an additional source
934 during that period contributing an average of 50% of OC probably due to use of wood for
935 heating of houses in winter and the festive period.

936 Lee et al. (2008a) used CMB and UNMIX receptor models to apportion sources of PM_{2.5}
937 aerosols collected between March 2001 and February 2001 in Korea. The CMB results
938 identified diesel vehicle exhaust as the major contributor to PM (33%), with meat cooking
939 contributing 12% of the PM_{2.5} mass measured. Other sources identified were secondary
940 sulphate (15%), secondary organic carbon (9%), urban dust, Asian dust, biomass burning, sea
941 salt, residual oil combustion, gasoline vehicle exhaust, automobile lead and unknown
942 components (Lee et al., 2008a). The UNMIX on the other hand only identified seven PM_{2.5}
943 sources and apportioned 30% of the mass to diesel vehicles, 17% to secondary sulphate, 15%
944 from biomass burning, secondary nitrate (13%), gasoline vehicle, secondary organic carbon
945 and Asian dust, but not cooking sources.

946 In Beijing (China), cooking was among the seven emission sources of particulate organic
947 matter identified (Wang et al., 2009). Like other studies, the other sources included gasoline
948 /diesel vehicles and vegetative burning in addition to coal burning in this case. The CMB
949 model established that contribution from cooking was actually higher during the summer,
950 whilst the biomass burning contribution was the highest during the winter (Wang et al.,
951 2009).

952 At a heavily polluted urban site in central California, molecular marker CMB was carried out
953 on ultrafine airborne particulate matter. Meat cooking was identified to account for 33-67%
954 of the PM_{0.1} at the urban site compared to diesel engines which accounted for 15-21%. At a
955 rural site meat cooking contributed 22-26% of the PM_{0.1} OC, and diesel engines accounted
956 for 8-9% (Ham and Kleeman, 2011). As regards the organic carbon of the larger PM_{1.8}
957 particles, meat cooking contributed less to the PM at the rural site than diesel engines; while
958 at the urban site the contribution from meat was still higher than from diesel engines. Lower
959 OC contributions were estimated compared to the measured concentrations, which implies an

960 unidentified contribution of either secondary organic aerosol (SOA) or oxidized primary
961 organic aerosol (POA). They estimated that meat cooking led to 0.01-0.025 $\mu\text{g}/\text{m}^3$ of $\text{PM}_{0.1}$
962 (Ham and Kleeman, 2011).

963 The above studies have illustrated the importance that cooking has as a source contributing to
964 organic aerosol. The studies have also identified that in order to reduce particle pollution,
965 especially in populated metropolitan areas, efforts should focus on controlling cooking as
966 well as other particle sources such as traffic emissions.

967

968 **5.2. Diagnostic Ratio and Ratio:Ratio Plots**

969 Diagnostic ratio is a binary ratio method for source identification which involves comparing
970 ratios of pairs of frequently found compounds emitted to distinguish between different known
971 sources. It is usually used with caution as a) it is often difficult to discriminate between some
972 sources; b) its interpretation depends on the ratio considered; and c) the profile chosen can
973 vary in the presence of highly reactive compounds and thus can introduce bias.

974 Similarly, plots of concentrations of different markers against each other has been a useful
975 tool to examine correlations among different markers. Robinson et al. (2006) made plots of
976 various ambient species, focussing on only five important markers for cooking; n-
977 hexadecanoic (palmitic) acid, n-octadecanoic (stearic) acid, 9-hexadecenoic (palmitoleic)
978 acid, 9-octadecenoic (oleic) acid, and cholesterol. Oleic and palmitoleic acid concentrations
979 as well as stearic and palmitic acid concentrations were well correlated with a slope of one
980 implying a single dominant source for the alkenoic acids. Only a slight correlation was
981 observed between cholesterol and palmitic acid but the scatter was comparable to
982 measurement uncertainty, so the sources could have been the same. Saturated and unsaturated

983 acids however, when plotted against one another showed no correlation (e.g. palmitic acid
984 against palmitoleic acid) (Robinson et al., 2006). Assuming they are chemically stable, it was
985 concluded that these acids have different dominant sources. Ratio–ratio plots aid in the
986 inference of potential source profiles. These plots should be examined using different ratio
987 species and different combinations of source specific markers to develop a good
988 understanding of ambient data and source profiles. Robinson et al. (2006) made use of such
989 plots of acids (two alkanonic and alkenoic) normalised by cholesterol and observed a good
990 correlation in ambient data by displaying well organised ratio to ratio plots.

991 Normalisation is a general approach to reduce the anomalies in large data sets. Scatters along
992 the diagonals of the plots can be attributed to measurement uncertainty or the variability of
993 emissions of species. The reference specie used to normalise the concentrations of the two
994 target species affects the exact organisation of data in a ratio-ratio plot. Changing the
995 reference specie should not alter the likely conclusion about the source profile combination
996 (Robinson et al., 2006) as illustrated in Figure 2. The change however causes the location of
997 both the source profiles and ambient data in the plots to shift. Cholesterol has been found to
998 be a good reference for food cooking markers and so has been generally used.

999 Diagnostic ratios for PAHs, such as BeP/(BeP + BaP), IND/(IND + BghiP), Cor/BeP and
1000 BghiP/BeP have been widely used in the investigation of their origins and to aid in the
1001 identification of the possible emission sources in air samples. See et al. (2006) used this
1002 technique in combination with other statistical methods. The ratios were calculated to provide
1003 insight into the origins as well as markers or tracers of pollution sources. The ratios of
1004 Phe/(ant + Phe) (structural isomers of molecular weight MW = 178), Flt/(Flt + Pyr) (MW =
1005 202), BaA/(BaA + Chr) (MW = 228) and Ind/(Ind + BPe) (MW = 276) were evaluated based
1006 on mean concentrations (See et al., 2006).

1007 ANT/(ANT + PHE) ratios were used by Gu et al. (2010) to indicate combustion sources
1008 while FLU/(FLU+PYR) ratio < 0.2 was taken as an indication of PAHs from petroleum
1009 sources, 0.4–0.5 for combustion of fuel and > 0.5 for grass combustion (Miguel and Pereira,
1010 1989; Gu et al., 2010).

1011 Table 19 shows some diagnostic ratios obtained for culinary techniques and from vehicle
1012 emissions as a comparison. The ratios from See et al. (2006) and See and Balasubramanian
1013 (2008) are relatively unaffected by type of cooking and there is some overlap in cooking
1014 ratios with those from traffic, making quantitative differentiation impossible.

1015 Some size resolved source apportionment studies have used molecular marker to organic
1016 carbon ratios as chemical signatures for source contribution identification and good tracers
1017 have been identified for molecular markers with similar size distribution for EC and OC
1018 (Kleeman et al., 2008). Kleeman et al. (2008) found that the size distribution for cholesterol
1019 was highly correlated ($R^2 > 0.9$) with both OC and EC size distributions further suggesting
1020 that cholesterol can serve as an appropriate tracer for meat cooking contributions, as
1021 suggested by previous authors (Robinson et al., 2006). The most abundant PAH measured in
1022 emissions from meat cooking was phenanthrene, with smaller concentrations of fluoranthene
1023 and pyrene. Higher ambient concentrations of these compounds were observed from other
1024 sources such as diesel engines (Kleeman et al, 2008).

1025 In Xiamen, China, diagnostic ratios and PCA were used to identify that cooking sources,
1026 gasoline and diesel vehicle exhaust, industry emissions and coal combustion were the sources
1027 of particle bound PAH (Hong et al., 2007).

1028 Ho et al. (2010) used the ratio of octadecanoic acid to hexadecanoic acid to identify cooking
1029 emissions as a source of pollution from a set of organic species measured during the air

1030 quality sampling campaign in Beijing in 2006. The ratio for cooking was between 0.39 and
1031 0.85 with an average value of 0.36 (Ho et al., 2010).

1032

1033 **5.3. Multivariate Statistical Methods**

1034 Principal Component Analysis is multivariate statistical technique which reduces the number
1035 of variables and groups into factor variables with similar characteristics. The set variables are
1036 transformed into a smaller set of linear combinations to analyse the variance of the original
1037 set. The selection of compounds and elements used in the model are based on their signal to
1038 noise ratio, their suitability for use as definite source markers and their percentage of values
1039 above the quantification limit (Brinkman et al., 2009).

1040 PCA has been found to assist effectively in identification of factors affecting personal
1041 exposure to PM using organic tracers by Brinkman et al. (2009). They measured a broad
1042 distribution of organic tracer compounds in PM personal samples and found that multiple
1043 organic marker species for the same source loaded the same factor, with meat cooking being
1044 observed to have high correlation with organic carbon concentrations ($r = 0.84$) (Brinkman et
1045 al., 2009).

1046 As most primary tracers are emitted from multiple sources, an assemblage of compounds as a
1047 group, rather than an individual tracer is used for indicating the quantity of cooking aerosol in
1048 the atmosphere. Cholesterol supplemented with monoglycerides, oleic acid and nonanal have
1049 been identified as potential primary tracers for meat cooking (Nolte et al., 1999).

1050 PCA was also used to analyse sources of PM₁₀ using concentrations of ambient air polycyclic
1051 aromatic hydrocarbons bound to PM at various sites consisting of a garden site, industrial-
1052 traffic intersection, a residential site and an island site in the city of Xiamen in China. In the

1053 summer period, five components were identified by the model, with 12% of the variance
1054 attributed to cooking fuels, which was indicated by high loadings of Chry and moderate
1055 loadings of BaA, Acen and An (Hong et al., 2007). The highest contributions were actually
1056 seen from gasoline and diesel vehicles exhausts throughout the study period.

1057 $PM_{2.5}$ measurements from some sites in California (N=23) were analysed using Unmix and
1058 PMF. Eight factors were obtained for both models, which included cooking, marine sea salt,
1059 fugitive dust, agriculture-dairy, secondary aerosol motor vehicle and residential wood
1060 combustion (Chen et al., 2007).

1061

1062 **6. CONCLUSION**

1063 Cooking has been shown to be a source of particulate matter both indoors and outdoors in
1064 cities around the world. Apportionment studies that have identified this source have made use
1065 of source profiles mainly derived in the United States as the earlier studies were carried out in
1066 that region.

1067 The composition of cooking aerosol is highly diverse, depending upon factors such as the raw
1068 food composition, cooking oil (if used), cooking temperature and cooking style. This
1069 diversity implies that identification and quantification of cooking aerosol in the atmosphere is
1070 very difficult unless the source characteristics at the measurement location are well known.

1071 While further knowledge of source-related chemical composition would be beneficial,
1072 quantitative source apportionment will remain imprecise and potentially inaccurate in
1073 situations with mixed cooking source types contributing to atmospheric concentrations. This
1074 provides a considerable challenge to Chemical Mass Balance models where it is necessary to
1075 input a source profile. It is less problematic for the now common studies in which PMF is

1076 used to disaggregate mass spectral data from AMS instruments. However, as yet such studies
1077 are unable to identify specific cooking styles and considerable skill is needed to achieve an
1078 optimal separation of the cooking organic aerosol (COA) factor from other factors
1079 contributing to organic carbon concentrations.

1080 There is a need for an enhanced understanding of cooking emissions around the world and of
1081 the effects upon human health. This review has found that Chinese cooking can lead to a
1082 much greater contribution of PAHs to particulate organic matter relative to western-style fast
1083 food cooking. The studies reviewed identified potential candidates of tracers for cooking
1084 emissions as tetradecanoic, hexadecanoic and octadecanoic acids, nonanal, lactones and
1085 levoglucosan. Western-style fast food cooking tracers were 9-octadecenoic, hexanedioic and
1086 nonanedioic acids; whilst Chinese cooking tracers were oleic acid, mannosan and galactosan.
1087 Further studies will go a long way to provide further insight to verify these choices and to
1088 determine whether these choices vary with geographical location and other culinary
1089 techniques.

1090 This review also shows that cooking aerosol comprises particles largely generated in the
1091 ultrafine size region. These particles have an associated large surface area and are capable of
1092 deposition in the respiratory system with high efficiency. Improved capture of cooking
1093 emissions above the stove by fume extraction and filtration would contribute substantially to
1094 improvements in both indoor and outdoor air quality, and hence a reduction in human
1095 exposure.

1096

References

- 1097
- 1098
- 1099 Abt, E., H. H. Suh, G. Allen and P. Koutrakis, 2000. Characterization of Indoor Particle Sources: A
1100 Study Conducted in the Metropolitan Boston Area. *Environmental Health Perspectives* 108,
1101 35-44.
- 1102 Aiken, A. C., B. de Foy, C. Wiedinmyer, P. F. DeCarlo, I. M. Ulbrich, M. N. Wehrli, S. Szidat, A. S. H.
1103 Prevot, J. Noda, L. Wacker, R. Volkamer, E. Fortner, J. Wang, A. Laskin, V. Shutthanandan, J.
1104 Zheng, R. Zhang, G. Paredes-Miranda, W. P. Arnott, L. T. Molina, G. Sosa, X. Querol and J. L.
1105 Jimenez, 2010. Mexico city aerosol analysis during MILAGRO using high resolution aerosol
1106 mass spectrometry at the urban supersite (T0) – Part 2: Analysis of the biomass burning
1107 contribution and the non-fossil carbon fraction. *Atmos. Chem. Phys.* 10, 5315-5341.
- 1108 Allan, J. D., A. E. Delia, H. Coe, K. N. Bower, M. R. Alfarra, J. L. Jimenez, A. M. Middlebrook, F.
1109 Drewnick, T. B. Onasch, M. R. Canagaratna, J. T. Jayne and D. R. Worsnop, 2004. A
1110 generalised method for the extraction of chemically resolved mass spectra from Aerodyne
1111 aerosol mass spectrometer data. *Journal of Aerosol Science* 35, 909-922.
- 1112 Allan, J. D., P. I. Williams, W. T. Morgan, C. L. Martin, M. J. Flynn, J. Lee, E. Nemitz, G. J. Phillips, M.
1113 W. Gallagher and H. Coe, 2010. Contributions from transport, solid fuel burning and cooking
1114 to primary organic aerosols in two UK cities. *Atmos. Chem. Phys.* 10, 647-668.
- 1115 Allen, R., L. Wallace, T. Larson, L. Sheppard and L. J. S. Liu, 2004. Estimated hourly personal
1116 exposures to ambient and nonambient particulate matter among sensitive populations in
1117 Seattle, Washington. *Journal of the Air & Waste Management Association* 54, 1197-1211.
- 1118 Baltensperger, U. and A. Prévôt, 2008. Chemical analysis of atmospheric aerosols. *Analytical and*
1119 *Bioanalytical Chemistry* 390, 277-280.
- 1120 Barham, P., 1950. *The Science of Cooking*, Springer-Verlag Berlin and Heidelberg GmbH & Co. K.
- 1121 Birch, M. E. and R. A. Cary, 1996. Elemental carbon-based method for monitoring occupational
1122 exposures to particulate diesel exhaust. *Aerosol Science and Technology* 25, 221-241.
- 1123 Brinkman, G. L., J. B. Milford, J. J. Schauer, M. M. Shafer and M. P. Hannigan, 2009. Source
1124 identification of personal exposure to fine particulate matter using organic tracers.
1125 *Atmospheric Environment* 43, 1972-1981.
- 1126 Brunekreef, B., N. A. H. Janssen, M. Oldenwening, K. Meliefste, G. Hoek, T. Lanki, K. L. Timonen, M.
1127 Vallius, J. Pekkanen and R. Van Grieken, 2005. Personal, indoor and outdoor exposures of
1128 PM_{2.5} and its components for groups of cardiovascular patients in Amsterdam and
1129 Helsinki. Research Report 127. Boston MA, Health Effects Institute.
- 1130 Buonanno, G., G. Johnson, L. Morawska and L. Stabile, 2011. Volatility Characterization of Cooking-
1131 Generated Aerosol Particles. *Aerosol Science and Technology* 45, 1069-1077.
- 1132 Buonanno, G., L. Morawska and L. Stabile, 2009. Particle emission factors during cooking activities.
1133 *Atmospheric Environment* 43, 3235-3242.
- 1134 Buonanno, G., L. Morawska, L. Stabile and A. Viola, 2010. Exposure to particle number, surface area
1135 and PM concentrations in pizzerias. *Atmospheric Environment* 44, 3963-3969.
- 1136 Cabada, J. C., S. N. Pandis and A. L. Robinson, 2002. Sources of atmospheric carbonaceous
1137 particulate matter in Pittsburgh, Pennsylvania. *Journal of the Air & Waste Management*
1138 *Association* 52, 732-741.
- 1139 Canagaratna, M. R., J. T. Jayne, J. L. Jimenez, J. D. Allan, M. R. Alfarra, Q. Zhang, T. B. Onasch, F.
1140 Drewnick, H. Coe, A. Middlebrook, A. Delia, L. R. Williams, A. M. Trimborn, M. J. Northway, P.
1141 F. DeCarlo, C. E. Kolb, P. Davidovits and D. R. Worsnop, 2007. Chemical and microphysical
1142 characterization of ambient aerosols with the aerodyne aerosol mass spectrometer. *Mass*
1143 *Spectrom Rev* 26, 185-222.
- 1144 Chao, C. Y. and E. C. Cheng, 2002. Source Apportionment of Indoor PM_{2.5} and PM₁₀ in Homes.
1145 *Indoor and Built Environment* 11, 27-37.
- 1146 Chappell, W. R., C. O. Abernathy, R. L. Calderon and D. J. Thomas, Eds, 2003. Arsenic Exposure and
1147 Health Effects V. Proceedings of the Fifth International Conference on Arsenic Exposure and

1148 Health Effects, July 14–18, 2002. Fifth International Conference on Arsenic Exposure and
1149 Health Effects. San Diego, California.

1150 Chen, B. H. and Y. C. Chen, 2001. Formation of polycyclic aromatic hydrocarbons in the smoke from
1151 heated model lipids and food lipids. *Journal Of Agricultural And Food Chemistry* 49, 5238-
1152 5243.

1153 Chen, L. W. A., J. G. Watson, J. C. Chow and K. L. Magliano, 2007. Quantifying PM_{2.5} source
1154 contributions for the San Joaquin Valley with multivariate receptor models. *Environmental*
1155 *science & technology* 41, 2818-2826.

1156 Chen, Y. C. and B. H. Chen, 2003. Determination of Polycyclic Aromatic Hydrocarbons in Fumes from
1157 Fried Chicken Legs. *Journal of Agricultural and Food Chemistry* 51, 4162-4167.

1158 Chiang, T. A., P. F. Wu and Y. C. Ko, 1999. Identification of carcinogens in cooking oil fumes. *Environ*
1159 *Res* 81, 18-22.

1160 Chow, J. C. and J. G. Watson, 2002. Review of PM_{2.5} and PM₁₀ Apportionment for Fossil Fuel
1161 Combustion and Other Sources by the Chemical Mass Balance Receptor Model. *Energy &*
1162 *Fuels* 16, 222-260.

1163 Chow, J. C., J. G. Watson, H. Kuhns, V. Etyemezian, D. H. Lowenthal, D. Crow, S. D. Kohl, J. P.
1164 Engelbrecht and M. C. Green, 2004. Source profiles for industrial, mobile, and area sources
1165 in the Big Bend Regional Aerosol Visibility and Observational study. *Chemosphere* 54, 185-
1166 208.

1167 Clougherty, J. E., E. A. Houseman and J. I. Levy, 2011. Source apportionment of indoor residential
1168 fine particulate matter using land use regression and constrained factor analysis. *Indoor Air*
1169 21, 53-66.

1170 Delgado-Saborit, J. M., N. J. Aquilina, C. Meddings, S. Baker and R. M. Harrison, 2011. Relationship of
1171 personal exposure to volatile organic compounds to home, work and fixed site outdoor
1172 concentrations. *Science of the Total Environment* 409, 478-488.

1173 Dennekamp, M., S. Howarth, C. A. J. Dick, J. W. Cherrie, K. Donaldson and A. Seaton, 2001. Ultrafine
1174 particles and nitrogen oxides generated by gas and electric cooking. *Occupational and*
1175 *Environmental Medicine* 58, 511-516.

1176 Diapouli, E., K. Eleftheriadis, A. A. Karanasiou, S. Vratolis, O. Hermansen, I. Colbeck and M. Lazaridis,
1177 2011. Indoor and Outdoor Particle Number and Mass Concentrations in Athens. Sources,
1178 Sinks and Variability of Aerosol Parameters. *Aerosol Air Qual. Res.* 11, 632-642.

1179 Fang, G. C., C. N. Chang, Y. S. Wu, P. P. C. Fu, D. G. Yang and C. C. Chu, 1999. Characterization of
1180 chemical species in PM_{2.5} and PM₁₀ aerosols in suburban and rural sites of central Taiwan.
1181 *Science of the Total Environment* 234, 203-212.

1182 Figueroa, D. A., C. J. Rodriguez-Sierra and B. D. Jimenez-Velez, 2006. Concentrations of Ni and V,
1183 other heavy metals, arsenic, elemental and organic carbon in atmospheric fine particles
1184 (PM_{2.5}) from Puerto Rico. *Toxicology And Industrial Health* 22, 87-99.

1185 Fortmann, R., P. Kariher and C. Clayton, 2001. Indoor Air Quality: Residential Cooking Exposures,
1186 California Air Resources Board.

1187 Fraser, M. P., G. R. Cass, B. R. T. Simoneit and R. A. Rasmussen, 1997. Air quality model evaluation
1188 data for organics .4. C-2-C-36 non-aromatic hydrocarbons. *Environmental Science &*
1189 *Technology* 31, 2356-2367.

1190 Fraser, M. P., K. Lakshmanan, S. G. Fritz and B. Ubanwa, 2002. Variation in composition of fine
1191 particulate emissions from heavy-duty diesel vehicles. *J. Geophys. Res.* 107, 8346.

1192 Fraser, M. P., Z. W. Yue and B. Buzcu, 2003. Source apportionment of fine particulate matter in
1193 Houston, TX, using organic molecular markers. *Atmospheric Environment* 37, 2117-2123.

1194 Glytsos, T., J. Ondracek, L. Dzumbova, I. Kopanakis and M. Lazaridis, 2010. Characterization of
1195 particulate matter concentrations during controlled indoor activities. *Atmospheric*
1196 *Environment* 44, 1539-1549.

1197 Goldstein, A. H. and I. E. Galbally, 2007. Known and Unexplored Organic Constituents in the Earth's
1198 Atmosphere. *Environmental science & technology* 41, 1514-1521.

- 1199 Grosjean, E., D. Grosjean, M. P. Fraser and G. R. Cass, 1996. Air quality model evaluation data for
1200 organics .2. C-1-C-14 carbonyls in Los Angeles air. *Environmental Science & Technology* 30,
1201 2687-2703.
- 1202 Gu, Z., J. Feng, W. Han, L. Li, M. Wu, J. Fu and G. Sheng, 2010. Diurnal variations of polycyclic
1203 aromatic hydrocarbons associated with PM_{2.5} in Shanghai, China. *Journal of Environmental*
1204 *Sciences* 22, 389-396.
- 1205 Gundel, L. A., V. C. Lee, K. R. R. Mahanama, R. K. Stevens and J. M. Daisey, 1995. Direct
1206 determination of the phase distributions of semi-volatile polycyclic aromatic hydrocarbons
1207 using annular denuders. *Atmospheric Environment* 29, 1719.
- 1208 Ham, W. A. and M. J. Kleeman, 2011. Size-resolved source apportionment of carbonaceous
1209 particulate matter in urban and rural sites in central California. *Atmospheric Environment*
1210 45, 3988-3995.
- 1211 Harrison, R. and J. Yin, 2005. Characterisation of Particulate Matter in the United Kingdom.
1212 (Literature Review for Defra) Report for Defra, the National Assembly for Wales, the DoENI
1213 and the Scottish Executive.
- 1214 He, C., L. Morawska, J. Hitchins and D. Gilbert, 2004a. Contribution from indoor sources to particle
1215 number and mass concentrations in residential houses. *Atmospheric Environment* 38, 3405-
1216 3415.
- 1217 He, L.-Y., X.-F. Huang, L. Xue, M. Hu, Y. Lin, J. Zheng, R. Zhang and Y.-H. Zhang, 2011. Submicron
1218 aerosol analysis and organic source apportionment in an urban atmosphere in Pearl River
1219 Delta of China using high-resolution aerosol mass spectrometry. *J. Geophys. Res.* 116,
1220 D12304.
- 1221 He, L. Y., M. Hu, X. F. Huang, B. D. Yu, Y. H. Zhang and D. Q. Liu, 2004b. Measurement of emissions of
1222 fine particulate organic matter from Chinese cooking. *Atmospheric Environment* 38, 6557-
1223 6564.
- 1224 He, L. Y., M. Hu, L. Wang, X. F. Huang and Y. H. Zhang, 2004c. Characterization of fine organic
1225 particulate matter from Chinese cooking. *Journal Of Environmental Sciences-China* 16, 570-
1226 575.
- 1227 He, L. Y., Y. Lin, X. F. Huang, S. Guo, L. Xue, Q. Su, M. Hu, S. J. Luan and Y. H. Zhang, 2010.
1228 Characterization of high-resolution aerosol mass spectra of primary organic aerosol
1229 emissions from Chinese cooking and biomass burning. *Atmospheric Chemistry and Physics*
1230 10, 11535-11543.
- 1231 Hetland, R. B., M. Refsnes, T. Myran, B. V. Johansen, N. Uthus and P. E. Schwarze, 2000. Mineral
1232 and/or metal content as critical determinants of particle-induced release of IL-6 and IL-8
1233 from A549 cells. *Journal of Toxicology and Environmental Health-Part A* 60, 47-65.
- 1234 Hildemann, L. M., G. R. Markowski and G. R. Cass, 1991a. Chemical composition of emissions from
1235 urban sources of fine organic aerosol. *Environmental science & technology* 25, 744-759.
- 1236 Hildemann, L. M., M. A. Mazurek, G. R. Cass and B. R. T. Simonelt, 1991b. Quantitative
1237 characterization of urban sources of organic aerosol by high-resolution gas chromatography.
1238 *Environmental Science and Technology* 25, 1311-1325.
- 1239 Ho, K. F., S. C. Lee, S. S. H. Ho, K. Kawamura, E. Tachibana, Y. Cheng and T. Zhu, 2010. Dicarboxylic
1240 acids, ketocarboxylic acids, α-dicarbonyls, fatty acids, and benzoic acid in urban
1241 aerosols collected during the 2006 Campaign of Air Quality Research in Beijing (CAREBeijing-
1242 2006). *J. Geophys. Res.* 115, D19312.
- 1243 Ho, K. F., S. C. Lee, J. C. Yu, S. C. Zou and K. Fung, 2002. Carbonaceous characteristics of atmospheric
1244 particulate matter in Hong Kong. *Science of the Total Environment* 300, 59-67.
- 1245 Hong, H., H. Yin, X. Wang and C. Ye, 2007. Seasonal variation of PM₁₀-bound PAHs in the
1246 atmosphere of Xiamen, China. *Atmospheric Research* 85, 429-441.
- 1247 Hopke, P. K., Z. Ramadan, P. Paatero, G. A. Norris, M. S. Landis, R. W. Williams and C. W. Lewis, 2003.
1248 Receptor modeling of ambient and personal exposure samples: 1998 Baltimore Particulate
1249 Matter Epidemiology-Exposure Study. *Atmospheric Environment* 37, 3289-3302.

1250 Hsieh, L. T., Y. F. Wang, P. Li and K. C. Chen, 2011. Removal of Particle-bound Water-soluble Ions
1251 from Cooking Fume Using Bio-solution Wet Scrubber. . *Aerosol Air Qual. Res.* 11, 508–518.

1252 Huang, X. F., L. Y. He, M. Hu, M. R. Canagaratna, Y. Sun, Q. Zhang, T. Zhu, L. Xue, L. W. Zeng, X. G. Liu,
1253 Y. H. Zhang, J. T. Jayne, N. L. Ng and D. R. Worsnop, 2010. Highly time-resolved chemical
1254 characterization of atmospheric submicron particles during 2008 Beijing Olympic Games
1255 using an Aerodyne High-Resolution Aerosol Mass Spectrometer. *Atmos. Chem. Phys.*
1256 *Discuss.* 10, 13219-13251.

1257 Huboyo, H. S., S. Tohno and R. Q. Cao, 2011. Indoor PM_{2.5} Characteristics and CO Concentration
1258 Related to Water-Based and Oil-Based Cooking Emissions Using a Gas Stove. *Aerosol and Air*
1259 *Quality Research* 11, 401-411.

1260 Huffman, J. A., K. S. Docherty, C. Mohr, M. J. Cubison, I. M. Ulbrich, P. J. Ziemann, T. B. Onasch and J.
1261 L. Jimenez, 2009. Chemically-Resolved Volatility Measurements of Organic Aerosol from
1262 Different Sources. *Environmental Science & Technology* 43, 5351-5357.

1263 Hussein, T., T. Glytsos, J. Ondracek, P. Dohanyosova, V. Zdimal, K. Hameri, M. Lazaridis, J. Smolik and
1264 M. Kulmala, 2006. Particle size characterization and emission rates during indoor activities in
1265 a house. *Atmospheric Environment* 40, 4285-4307.

1266 IARC, 1994. Some industrial chemicals. . IARC Monographs on the Evaluation of Carcinogenic Risks
1267 to Humans Lyon, France, International Agency for Research on Cancer
1268 60, 1–560.

1269 IARC, 1995. Acrolein (Group 3). International Agency for Research on Cancer (IARC) - Summaries &
1270 Evaluations. Lyon, France, International Agency for Research on Cancer
1271 63, 337.

1272 IARC, 1999. Re-evaluation of some organic chemicals, hydrazine and hydrogen peroxide. IARC
1273 Monographs on the Evaluation of Carcinogenic Risks to Humans Lyon, France, International
1274 Agency for Research on Cancer
1275 71, 1–315. .

1276 IARC, 2006. Formaldehyde, 2-butoxyethanol and 1-tert-butoxypropan-2-ol. IARC Monographs on the
1277 Evaluation of Carcinogenic Risks to Humans Lyon, France, International Agency for Research
1278 on Cancer
1279 88, 1–478. .

1280 IARC, 2010. Household Use of Solid Fuels and High-temperature Frying. IARC Monographs on the
1281 Evaluation of Carcinogenic Risks to Humans Lyon, France, International Agency for Research
1282 on Cancer
1283 95.

1284 Kamens, R., C.-t. Lee, R. Wiener and D. Leith, 1991. A study of characterize indoor particles in three
1285 non-smoking homes. *Atmospheric Environment. Part A. General Topics* 25, 939-948.

1286 Kanakidou, M., J. H. Seinfeld, S. N. Pandis, I. Barnes, F. J. Dentener, M. C. Facchini, R. Van Dingenen,
1287 B. Ervens, A. Nenes, C. J. Nielsen, E. Swietlicki, J. P. Putaud, Y. Balkanski, S. Fuzzi, J. Horth, G.
1288 K. Moortgat, R. Winterhalter, C. E. L. Myhre, K. Tsigaridis, E. Vignati, E. G. Stephanou and J.
1289 Wilson, 2005. Organic aerosol and global climate modelling: a review. *Atmos. Chem. Phys.* 5,
1290 1053-1123.

1291 Kim, K. H., S. K. Pandey, E. Kabir, J. Susaya and R. J. C. Brown, 2011. The modern paradox of
1292 unregulated cooking activities and indoor air quality. *Journal of Hazardous Materials* 195, 1-
1293 10.

1294 Kleeman, M. J., M. A. Robert, S. G. Riddle, P. M. Fine, M. D. Hays, J. J. Schauer and M. P. Hannigan,
1295 2008. Size distribution of trace organic species emitted from biomass combustion and meat
1296 charbroiling. *Atmospheric Environment* 42, 3059-3075.

1297 Kleeman, M. J., J. J. Schauer and G. R. Cass, 1999. Size and Composition Distribution of Fine
1298 Particulate Matter Emitted from Wood Burning, Meat Charbroiling, and Cigarettes.
1299 *Environmental science & technology* 33, 3516-3523.

- 1300 Koistinen K.J., H. O., Rotko T., Edwards R.D., Moschandreas D., Jantunen M.J., 2001. Behavioral and
 1301 environmental determinants of personal exposures to PM_{2.5} in EXPOLIS } Helsinki, Finland.
 1302 Atmospheric Environment 35, 2473 - 2481.
- 1303 Lai, A. C. K. and Y. W. Ho, 2008. Spatial concentration variation of cooking-emitted particles in a
 1304 residential kitchen. Building and Environment 43, 871-876.
- 1305 Lai, S. C., K. F. Ho, Y. Y. Zhang, S. C. Lee, Y. Huang and S. C. Zou, 2010. Characteristics of Residential
 1306 Indoor Carbonaceous Aerosols: A Case Study in Guangzhou, Pearl River Delta Region.
 1307 Aerosol and Air Quality Research 10, 472-478.
- 1308 Lanz, V. A., M. R. Alfara, U. Baltensperger, B. Buchmann, C. Hueglin and A. S. H. Prevot, 2007. Source
 1309 apportionment of submicron organic aerosols at an urban site by factor analytical modelling
 1310 of aerosol mass spectra. Atmospheric Chemistry And Physics 7, 1503-1522.
- 1311 Lazaridis, M., V. Aleksandropoulou, J. Smolík, J. E. Hansen, T. Glytsos, N. Kalogerakis and E. Dahlin,
 1312 2006. Physico-chemical characterization of indoor/outdoor particulate matter in two
 1313 residential houses in Oslo, Norway: measurements overview and physical properties –
 1314 URBAN-AEROSOL Project. Indoor Air 16, 282-295.
- 1315 Lee, H., S. S. Park, K. W. Kim and Y. J. Kim, 2008a. Source identification of PM_{2.5} particles measured
 1316 in Gwangju, Korea. Atmospheric Research 88, 199-211.
- 1317 Lee, S., W. Liu, Y. Wang, A. G. Russell and E. S. Edgerton, 2008b. Source apportionment of PM_{2.5}:
 1318 Comparing PMF and CMB results for four ambient monitoring sites in the southeastern
 1319 United States. Atmospheric Environment 42, 4126-4137.
- 1320 Lee, S. C., K. F. Ho, L. Y. Chan, B. Zielinska and J. C. Chow, 2001. Polycyclic aromatic hydrocarbons
 1321 (PAHs) and carbonyl compounds in urban atmosphere of Hong Kong. Atmospheric
 1322 Environment 35, 5949-5960.
- 1323 Levy, J. I., T. Dumyahn and J. D. Spengler, 2002. Particulate matter and polycyclic aromatic
 1324 hydrocarbon concentrations in indoor and outdoor microenvironments in Boston,
 1325 Massachusetts. Journal Of Exposure Analysis And Environmental Epidemiology 12, 104-114.
- 1326 Li, C. S., W. H. Lin and F. T. Jenq, 1993. SIZE DISTRIBUTIONS OF SUBMICROMETER AEROSOLS FROM
 1327 COOKING. Environment International 19, 147-154.
- 1328 Li, C. T., Y. C. Lin, W. J. Lee and P. J. Tsai, 2003. Emission of Polycyclic Aromatic Hydrocarbons and
 1329 Their Carcinogenic Potencies from Cooking Sources to the Urban Atmosphere.
 1330 Environmental Health Perspectives 111, 483-487.
- 1331 Lin, L., M. L. Lee and D. J. Eatough, 2010. Review of Recent Advances in Detection of Organic Markers
 1332 in Fine Particulate Matter and Their Use for Source Apportionment. Journal of the Air &
 1333 Waste Management Association 60, 3-25.
- 1334 Long, C. M., H. H. Suh and P. Koutrakis, 2000. Characterization of indoor particle sources using
 1335 continuous mass and size monitors. J Air Waste Manag Assoc 50, 1236-1250.
- 1336 Marcazzan, G. M., S. Vaccaro, G. Valli and R. Vecchi, 2001. Characterisation of PM₁₀ and PM_{2.5}
 1337 particulate matter in the ambient air of Milan (Italy). Atmospheric Environment 35, 4639-
 1338 4650.
- 1339 Massey, D., A. Kulshrestha, J. Masih and A. Taneja, 2012. Seasonal trends of PM₁₀, PM_{5.0}, PM_{2.5} &
 1340 PM_{1.0} in indoor and outdoor environments of residential homes located in North-Central
 1341 India. Building and Environment 47, 223-231.
- 1342 McDonald, J. D., B. Zielinska, E. M. Fujita, J. C. Sagebiel, J. C. Chow and J. G. Watson, 2003. Emissions
 1343 from charbroiling and grilling of chicken and beef. Journal of the Air & Waste Management
 1344 Association 53, 185-194.
- 1345 Miguel, A. H. and P. A. P. Pereira, 1989. Benzo(k)fluoranthene, Benzo(ghi)perylene, and
 1346 Indeno(1,2,3-cd)pyrene: New Tracers of Automotive Emissions in Receptor Modeling.
 1347 Aerosol Science and Technology 10, 292-295.
- 1348 Millar, G., T. Abel, J. Allen, P. Barn, M. Noullett, J. Spagnol and P. L. Jackson, 2010. Evaluating Human
 1349 Exposure to Fine Particulate Matter Part II: Modeling. Geography Compass 4, 731-749.

1350 Mohr, C., P. F. DeCarlo, M. F. Heringa, R. Chirico, J. G. Slowik, R. Richter, C. Reche, A. Alastuey, X.
1351 Querol, R. Seco, J. Peñuelas, J. L. Jiménez, M. Crippa, R. Zimmermann, U. Baltensperger and
1352 A. S. H. Prévôt, 2011. Identification and quantification of organic aerosol from cooking and
1353 other sources in Barcelona using aerosol mass spectrometer data. *Atmos. Chem. Phys.*
1354 *Discuss.* 11, 27383-27420.

1355 Mohr, C., J. A. Huffman, M. J. Cubison, A. C. Aiken, K. S. Docherty, J. R. Kimmel, I. M. Ulbrich, M.
1356 Hannigan and J. L. Jimenez, 2009. Characterization of Primary Organic Aerosol Emissions
1357 from Meat Cooking, Trash Burning, and Motor Vehicles with High-Resolution Aerosol Mass
1358 Spectrometry and Comparison with Ambient and Chamber Observations. *Environmental*
1359 *science & technology* 43, 2443-2449.

1360 Nolte, C. G., J. J. Schauer, G. R. Cass and B. R. T. Simoneit, 1999. Highly Polar Organic Compounds
1361 Present in Meat Smoke. *Environmental science & technology* 33, 3313-3316.

1362 Ozkaynak, H., J. Xue, J. Spengler, L. Wallace, E. Pellizzari and P. Jenkins, 1996. Personal exposure to
1363 airborne particles and metals: Results from the particle team study in Riverside, California.
1364 *Journal of Exposure Analysis and Environmental Epidemiology* 6, 57-78.

1365 Pant, P. and R. M. Harrison, 2012. Critical review of receptor modelling for particulate matter: A case
1366 study of India. *Atmospheric Environment* 49, 1-12.

1367 Robinson, A. L., R. Subramanian, N. M. Donahue, A. Bernardo-Bricker and W. F. Rogge, 2006. Source
1368 apportionment of molecular markers and organic aerosol. 3. Food cooking emissions.
1369 *Environmental Science & Technology* 40, 7820-7827.

1370 Roe, S. M., M. D. Spivey, H. C. Lindquist, P. Hemmer and R. Huntley, 2005. National Emissions
1371 Inventory for Commercial Cooking. The U.S. Environmental Protection Agency (EPA)
1372 Emission Factor and Inventory Group (EFIG)

1373 Rogge, W. F., 2000. Investigating Indoor Air Quality by using a Dynamic and Modular Source Testing
1374 Approach for Indoor Cooking; Final Report to the Center for Indoor Air Research (CIAR).

1375 Rogge, W. F., L. M. Hildemann, M. A. Mazurek, G. R. Cass and B. R. T. Simoneit, 1991. Sources of fine
1376 organic aerosol. 1. Charbroilers and meat cooking operations. *Environmental science &*
1377 *technology* 25, 1112-1125.

1378 Rogge, W. F., L. M. Hildemann, M. A. Mazurek, G. R. Cass and B. R. T. Simoneit, 1993a. Sources of
1379 fine organic aerosol. 4. Particulate abrasion products from leaf surfaces of urban plants.
1380 *Environmental science & technology* 27, 2700-2711.

1381 Rogge, W. F., L. M. Hildemann, M. A. Mazurek, G. R. Cass and B. R. T. Simoneit, 1993b. Sources of
1382 fine organic aerosol. 5. Natural gas home appliances. *Environmental science & technology*
1383 27, 2736-2744.

1384 Rogge, W. F., L. M. Hildemann, M. A. Mazurek, G. R. Cass and B. R. T. Simoneit, 1997. Sources of Fine
1385 Organic Aerosol. 8. Boilers Burning No. 2 Distillate Fuel Oil. *Environmental science &*
1386 *technology* 31, 2731-2737.

1387 Scapellato, M. L., C. Canova, A. de Simone, M. Carrieri, P. Maestrelli, L. Simonato and G. B.
1388 Bartolucci, 2009. Personal PM(10) exposure in asthmatic adults in Padova, Italy: seasonal
1389 variability and factors affecting individual concentrations of particulate matter. *International*
1390 *Journal Of Hygiene And Environmental Health* 212, 626-636.

1391 Schauer, J. J. and G. R. Cass, 2000. Source Apportionment of Wintertime Gas-Phase and Particle-
1392 Phase Air Pollutants Using Organic Compounds as Tracers. *Environmental science &*
1393 *technology* 34, 1821-1832.

1394 Schauer, J. J., M. J. Kleeman, G. R. Cass and B. R. T. Simoneit, 1999a. Measurement of Emissions from
1395 Air Pollution Sources. 1. C1 through C29 Organic Compounds from Meat Charbroiling.
1396 *Environmental science & technology* 33, 1566-1577.

1397 Schauer, J. J., M. J. Kleeman, G. R. Cass and B. R. T. Simoneit, 1999b. Measurement of Emissions from
1398 Air Pollution Sources. 2. C1 through C30 Organic Compounds from Medium Duty Diesel
1399 Trucks. *Environmental science & technology* 33, 1578-1587.

1400 Schauer, J. J., M. J. Kleeman, G. R. Cass and B. R. T. Simoneit, 2002. Measurement of Emissions from
1401 Air Pollution Sources. 4. C1–C27 Organic Compounds from Cooking with Seed Oils.
1402 Environmental science & technology 36, 567-575.

1403 Schauer, J. J., W. F. Rogge, L. M. Hildemann, M. A. Mazurek, G. R. Cass and B. R. T. Simoneit, 1996.
1404 Source apportionment of airborne particulate matter using organic compounds as tracers.
1405 Atmospheric Environment 30, 3837-3855.

1406 See, S. W. and R. Balasubramanian, 2006a. Physical characteristics of ultrafine particles emitted
1407 from different gas cooking methods. Aerosol and Air Quality Research 6, 82-96.

1408 See, S. W. and R. Balasubramanian, 2006b. Risk assessment of exposure to indoor aerosols
1409 associated with Chinese cooking. Environmental Research 102, 197-204.

1410 See, S. W. and R. Balasubramanian, 2008. Chemical characteristics of fine particles emitted from
1411 different gas cooking methods. Atmospheric Environment 42, 8852-8862.

1412 See, S. W., S. Karthikeyana and R. Balasubramanian, 2006. Health risk assessment of occupational
1413 exposure to particulate-phase polycyclic aromatic hydrocarbons associated with Chinese,
1414 Malay and Indian cooking. Journal of Environmental Monitoring 8, 369-376.

1415 Seinfeld, J. H. and J. F. Pankow, 2003. Organic atmospheric particulate material. Annu Rev Phys
1416 Chem 54, 121-140.

1417 Siegmann, K. and K. Sattler, 1996. Aerosol from hot cooking oil, a possible health hazard. Journal Of
1418 Aerosol Science 27, Supplement 1, S493-S494.

1419 Sioutas, C., R. J. Delfino and M. Singh, 2005. Exposure assessment for atmospheric ultrafine particles
1420 (UFPs) and implications in epidemiologic research. Environ Health Perspect 113, 947-955.

1421 Sjaastad, A. K., R. B. Jorgensen and K. Svendsen, 2010. Exposure to polycyclic aromatic hydrocarbons
1422 (PAHs), mutagenic aldehydes and particulate matter during pan frying of beefsteak. Occup
1423 Environ Med 67, 228-232.

1424 Sjaastad, A. K. and K. Svendsen, 2008. Exposure to Mutagenic Aldehydes and Particulate Matter
1425 During Panfrying of Beefsteak with Margarine, Rapeseed Oil, Olive Oil or Soybean Oil. Annals
1426 of Occupational Hygiene 52, 739-745.

1427 Sjaastad, A. K. and K. Svendsen, 2009. Exposure to Polycyclic Aromatic Hydrocarbons (PAHs),
1428 Mutagenic Aldehydes, and Particulate Matter in Norwegian à la Carte Restaurants. Annals of
1429 Occupational Hygiene 53, 723-729.

1430 Sjaastad, A. K., K. Svendsen and R. B. Jorgensen, 2008. Sub-micrometer particles: Their level and how
1431 they spread after pan frying of beefsteak. Indoor and Built Environment 17, 230-236.

1432 Stadler, R. H., I. Blank, N. Varga, F. Robert, J. Hau, P. A. Guy, M. C. Robert and S. Riediker, 2002.
1433 Acrylamide from Maillard reaction products. Nature 419, 449-450.

1434 Subramanian, R., N. M. Donahue, A. Bernardo-Bricker, W. F. Rogge and A. L. Robinson, 2007. Insights
1435 into the primary-secondary and regional-local contributions to organic aerosol and PM2.5
1436 mass in Pittsburgh, Pennsylvania. Atmospheric Environment 41, 7414-7433.

1437 Sun, Y. L., Q. Zhang, J. J. Schwab, K. L. Demerjian, W. N. Chen, M. S. Bae, H. M. Hung, O. Hogrefe, B.
1438 Frank, O. V. Rattigan and Y. C. Lin, 2011. Characterization of the sources and processes of
1439 organic and inorganic aerosols in New York city with a high-resolution time-of-flight aerosol
1440 mass spectrometer. Atmos. Chem. Phys. 11, 1581-1602.

1441 Svendsen, K., H. N. Jensen, I. Sivertsen and K. Sjaastad, 2002. Exposure to cooking fumes in
1442 restaurant kitchens in Norway. Annals Of Occupational Hygiene 46, 395-400.

1443 Swami, K., C. D. Judd, J. Orsini, K. X. Yang and L. Husain, 2001. Microwave assisted digestion of
1444 atmospheric aerosol samples followed by inductively coupled plasma mass spectrometry
1445 determination of trace elements. Fresenius Journal Of Analytical Chemistry 369, 63-70.

1446 Tan, C. C. L., K. N. Finney, Q. Chen, N. V. Russell, V. N. Sharifi and J. Swithenbank, 2012. Experimental
1447 Investigation of Indoor Air Pollutants in Residential Buildings. Indoor and Built Environment.

1448 To, W. M. and L. L. Yeung, 2011. Effect of Fuels on Cooking Fume Emissions. Indoor and Built
1449 Environment 20, 555-563.

- 1450 To, W. M., L. L. Yeung and C. Y. H. Chao, 2000. Characterisation of Gas Phase Organic Emissions from
1451 Hot Cooking Oil in Commercial Kitchens. *Indoor and Built Environment* 9, 228-232.
- 1452 USEPA, 1999. Emissions from Street Vendor Cooking Devices (Charcoal Grilling). Washington, DC, US
1453 Environmental Protection Agency, 1-551.
- 1454 Venkataraman, C. and S. K. Friedlander, 1994. SOURCE RESOLUTION OF FINE PARTICULATE
1455 POLYCYCLIC AROMATIC-HYDROCARBONS USING A RECEPTOR MODEL MODIFIED FOR
1456 REACTIVITY. *Journal of the Air & Waste Management Association* 44, 1103-1108.
- 1457 Wallace, L., R. Williams, A. Rea and C. Croghan, 2006. Continuous weeklong measurements of
1458 personal exposures and indoor concentrations of fine particles for 37 health-impaired North
1459 Carolina residents for up to four seasons. *Atmospheric Environment* 40, 399-414.
- 1460 Wallace, L. A., 1989. The Total Exposure Assessment Methodology (Team) Study - An Analysis Of
1461 Exposures, Sources, And Risks Associated With 4 Volatile Organic-Chemicals. *Journal Of The*
1462 *American College Of Toxicology* 8, 883-895.
- 1463 Wallace, L. A., S. J. Emmerich and C. Howard-Reed, 2004. Source Strengths of Ultrafine and Fine
1464 Particles Due to Cooking with a Gas Stove. *Environmental science & technology* 38, 2304-
1465 2311.
- 1466 Wan, M.-P., C.-L. Wu, G.-N. Sze To, T.-C. Chan and C. Y. H. Chao, 2011. Ultrafine particles, and PM2.5
1467 generated from cooking in homes. *Atmospheric Environment* 45, 6141-6148.
- 1468 Wang, Q., M. Shao, Y. Zhang, Y. Wei, M. Hu and S. Guo, 2009. Source apportionment of fine organic
1469 aerosols in Beijing. *Atmos. Chem. Phys.* 9, 8573-8585.
- 1470 Watson, J., E. Fujita, J. C. Chow, B. Zielinska, L. Richards, W. Neff and D. Dietrich, 1998. Northern
1471 Front Range Air Quality Study Final Report Prepared for Colorado State University, Fort
1472 Collins, CO, and EPRI, Palo Alto, CA, by Desert Research Institute: Reno, NV.
- 1473 Williams, B. J., A. H. Goldstein, N. M. Kreisberg, S. V. Hering, D. R. Worsnop, I. M. Ulbrich, K. S.
1474 Docherty and J. L. Jimenez, 2010. Major components of atmospheric organic aerosol in
1475 southern California as determined by hourly measurements of source marker compounds.
1476 *Atmos. Chem. Phys.* 10, 11577-11603.
- 1477 Wu, P.-F., T.-A. Chiang, L.-F. Wang, C.-S. Chang and Y.-C. Ko, 1998. Nitro-polycyclic aromatic
1478 hydrocarbon contents of fumes from heated cooking oils and prevention of mutagenicity by
1479 catechin. *Mutation Research/Fundamental and Molecular Mechanisms of Mutagenesis* 403,
1480 29-34.
- 1481 Yeung, L. L. and W. M. To, 2008. Size Distributions of the Aerosols Emitted from Commercial Cooking
1482 Processes. *Indoor and Built Environment* 17, 220-229.
- 1483 Zhang, Q., J. Jimenez, M. Canagaratna, I. Ulbrich, N. Ng, D. Worsnop and Y. Sun, 2011. Understanding
1484 atmospheric organic aerosols via factor analysis of aerosol mass spectrometry: a review.
1485 *Analytical and Bioanalytical Chemistry*, 1-23.
- 1486 Zhang, Q., J. L. Jimenez, M. R. Canagaratna, J. D. Allan, H. Coe, I. Ulbrich, M. R. Alfarra, A. Takami, A.
1487 M. Middlebrook, Y. L. Sun, K. Dzepina, E. Dunlea, K. Docherty, P. F. DeCarlo, D. Salcedo, T.
1488 Onasch, J. T. Jayne, T. Miyoshi, A. Shimono, S. Hatakeyama, N. Takegawa, Y. Kondo, J.
1489 Schneider, F. Drewnick, S. Borrmann, S. Weimer, K. Demerjian, P. Williams, K. Bower, R.
1490 Bahreini, L. Cottrell, R. J. Griffin, J. Rautiainen, J. Y. Sun, Y. M. Zhang and D. R. Worsnop,
1491 2007. Ubiquity and dominance of oxygenated species in organic aerosols in
1492 anthropogenically-influenced Northern Hemisphere midlatitudes. *Geophys. Res. Lett.* 34,
1493 L13801.
- 1494 Zhang, Q. F., R. H. Gangupomu, D. Ramirez and Y. F. Zhu, 2010. Measurement of Ultrafine Particles
1495 and Other Air Pollutants Emitted by Cooking Activities. *International journal of*
1496 *environmental research and public health* 7, 1744-1759.
- 1497 Zhao, W., P. K. Hopke, G. Norris, R. Williams and P. Paatero, 2006. Source apportionment and
1498 analysis on ambient and personal exposure samples with a combined receptor model and an
1499 adaptive blank estimation strategy. *Atmospheric Environment* 40, 3788-3801.

1500 Zhao, W. X., P. K. Hopke, E. W. Gelfand and N. Rabinovitch, 2007a. Use of an expanded receptor
1501 model for personal exposure analysis in schoolchildren with asthma. *Atmospheric*
1502 *Environment* 41, 4084-4096.

1503 Zhao, Y., M. Hu, S. Slanina and Y. Zhang, 2007b. Chemical compositions of fine particulate organic
1504 matter emitted from Chinese cooking. *Environ Sci Technol* 41, 99-105.

1505 Zhao, Y., M. Hu, S. Slanina and Y. Zhang, 2007c. The molecular distribution of fine particulate organic
1506 matter emitted from Western-style fast food cooking. *Atmospheric Environment* 41, 8163-
1507 8171.

1508 Zheng, M., G. R. Cass, J. J. Schauer and E. S. Edgerton, 2002. Source apportionment of PM_{2.5} in the
1509 southeastern United States using solvent-extractable organic compounds as tracers.
1510 *Environmental Science & Technology* 36, 2361-2371.

1511 Zheng, M., L. Ke, E. S. Edgerton, J. J. Schauer, M. Dong and A. G. Russell, 2006. Spatial distribution of
1512 carbonaceous aerosol in the southeastern United States using molecular markers and
1513 carbon isotope data. *J. Geophys. Res.* 111, D10S06.

1514 Zhu, L. and J. Wang, 2003. Sources and patterns of polycyclic aromatic hydrocarbons pollution in
1515 kitchen air, China. *Chemosphere* 50, 611-618.
1516
1517

1518
1519

1520
1521
1522
1523
1524
1525
1526
1527
1528
1529
1530
1531
1532
1533
1534
1535
1536
1537
1538
1539
1540
1541
1542
1543
1544
1545
1546
1547
1548
1549
1550
1551
1552
1553
1554
1555
1556
1557
1558
1559
1560
1561
1562
1563
1564
1565
1566
1567
1568
1569

TABLE LEGENDS

- Table 1. General cultural styles of cooking. Common ingredients, oils and spices used
- Table 2. National emissions rate (tonnes/year) of criteria pollutants from commercial cooking in the USA (Roe et al., 2005) and for highway vehicles (Chappell et al., 2003)
- Table 3. Particle mass and number concentration measured in indoor environments close to cooking activities
- Table 4. Size distribution studies for cooking aerosols
- Table 5. Particle diameter mode (i.e. diameter representing highest particle number concentration) of particle number size fraction distribution from cooking activities
- Table 6. Sampling, extraction and analysis of emissions from cooking
- Table 7. Main identified cooking marker species in the literature
- Table 8. Concentrations of organic compounds from western-style fast food and from Chinese cooking (ng/mg of particulate organic matter) (Zhao et al., 2007b,c)
- Table 9. Polycyclic aromatic hydrocarbons emitted from cooking food – indoor concentrations and emission factors (see columns for units)
- Table 10. Polycyclic aromatic hydrocarbons in fumes from heated oils - indoor concentrations and emission factors (see columns for units) - and in uncooked oil
- Table 11. N-alkanes emitted from cooking food - indoor concentrations and emission factors (see columns for units)
- Table 12. Fatty acids emitted from cooking food - indoor concentrations and emission factors (see columns for units)
- Table 13. Dicarboxylic acids emitted from cooking food - indoor concentrations and emission factors (see columns for units)
- Table 14. Aldehydes emitted from cooking food - indoor concentrations and emission factors (see columns for units)
- Table 15. Ketones and lactones emitted from cooking food - indoor concentrations and emission factors (see columns for units)
- Table 16. Concentrations of organic ions emitted from cooking food

1570 Table 17. Molecular markers, amides and other organic compounds emitted from
1571 cooking food - indoor concentrations and emission factors (see columns for
1572 units)

1573
1574 Table 18. EC/OC, metals and inorganic ions emitted from cooking food - indoor
1575 concentrations and emission factors (see columns for units)

1576
1577 Table 19. Comparison of diagnostic ratios of PAHs

1578

1579

1580

1581 **FIGURE LEGENDS**

1582

1583 Figure 1. Break down products of triglycerides (Nolte et al., 1999)

1584

1585 Figure 2. Marker to OC ratio for meat cooking profiles (Robinson et al., 2006)

1586

1587
1588
1589

Table 1. General cultural styles of cooking. Common ingredients, oils and spices used

COOKING STYLE	METHOD	INGREDIENTS	OIL	SPICES
Chinese	Stir fry, simmer, steam roast stew	MAIN- Meat type- Pork, sea food, poultry, beef, Vegetable-cabbage, carrots, cucumber, broccoli OTHERS- Eggs, ginger, hot pepper, scallion, garlic, rice, flour, peanuts, fruits	Soy beans Peanut oil Canola oil	essence of chicken, salt, peanut oil, light soy source, sugar
Western	Grill, broil, roast, deep fry, stew,	MAIN -Meat type-beef, chicken Vegetables-carrots, broccoli, OTHERS - milk, flour	Corn oil, vegetable oil olive oil,	Salt, black pepper, garlic, basil, parsley
Fast Food	Deep fry, stew	MAIN - Meat type- beef, chicken, Potatoes	Vegetable, butter, corn oil	Salt
African	Deep fry, boiling, stew	MAIN- Meat- beef, chicken, fish Vegetables-spinach OTHERS- yam, rice, plantain, banana.	Ground nut oil, palm oil, vegetable oil	Thyme, curry
Indian	Deep fry, boiling, stew	MAIN- Meat- fish and chicken OTHERS- rice, flour, beans, lentils, pearl millet, wheat flour, milk, yoghurt, plantain.	Vegetable oil, peanut oil, mustard oil, coconut oil, sesame oil,	Chilli pepper, black pepper, mustard seed, cumin, turmeric, ginger, cardamom, cinnamon, clove, garam masala, coriander, garlic, mustard seeds, nutmeg, mint
Malay	Deep fry, boiling, stew	MAIN- Meat-Fish, squids, prawns, crabs , chicken, beef and mutton. OTHERS-rice, noodles, yoghurt, coconut milk.	Vegetable oil, coconut oil, sesame oil,	Lemongrass, shallots, ginger, chillies, garlic, turmeric, lime leaves, laksa leaves, wild ginger flower buds or torch ginger and screwpine leaves, fennel, cumin, coriander, cardamom, cloves, star anise, mustard seeds, and nutmeg

1590

1591

1592

1593

1594

1595

1596 **Table 2. National emissions rate (tonnes/year) of criteria pollutants from commercial**
1597 **cooking in the USA (Roe et al., 2005) and for highway vehicles (Chappell et**
1598 **al., 2003)**

1599

1600

Pollutant	Total charbroiling	Deep frying	Flat griddle frying	Clamshell griddle frying	Under-fired charbroiling	Conveyorized charbroiling	Highway vehicles
VOC	115	1,170	39	940	7,200	2,100	4,400,000
CO	33,000		1,900		23,700	7,400	48,400,000
PM _{2.5}	79,300		11,900	910	58,300	8,200	135,000
PM ₁₀	85,500		15,700	1,100	60,300	8,500	192,000
PAH total	206		41		122	43	

1601

Table 3. Particle mass and number concentration measured in indoor environments close to cooking activities

Reference	Location	Comment	Concentration ($\mu\text{g}/\text{m}^3$)	Particle Number concentration (part/cm^3)
Li et al., 1993	Taiwan	Chicken		$1.2\text{-}2.6 \times 10^5$
Siegmann and Sattler, 1996	Switzerland	Rapeseed Oil		$2.5\text{-}4.5 \times 10^5$
Abt et al. 2000	US	Frying - $\text{PM}_{0.02\text{-}0.5}$	29	
		Frying - $\text{PM}_{0.7\text{-}10}$	19	
		Barbequing - $\text{PM}_{0.02\text{-}0.5}$	57	
		Barbequing - $\text{PM}_{0.7\text{-}10}$	12	
		Oven cooking - $\text{PM}_{0.02\text{-}0.5}$	50	
		Oven cooking - $\text{PM}_{0.7\text{-}10}$	8	
		Sauteing - $\text{PM}_{0.02\text{-}0.5}$	42	
		Sauteing - $\text{PM}_{0.7\text{-}10}$	294	
		Toasting - $\text{PM}_{0.02\text{-}0.5}$	45	
		Toasting - $\text{PM}_{0.7\text{-}10}$	8	
Dennekamp et al., 2001	UK	Frying vegetables (500 g) – gas stove		1.4×10^5
		Frying bacon (4 racers) – gas stove		5.9×10^5
		Frying vegetables (500 g) – electric stove		0.11×10^5
		Frying bacon (4 racers) – electric stove		1.6×10^5
		Bake cake – gas oven		0.9×10^5
		Bake cake – electric oven		0.3×10^5
		Roast meat and potatoes – gas oven		1.2×10^5
		Roast meat and potatoes – electric oven		0.2×10^5
		Toast – gas grill		1.4×10^5
		Toast – electric grill		1.4×10^5
Lee et al. 2001	China	$\text{PM}_{2.5}$ Chinese hot pot restaurant	81	
		$\text{PM}_{2.5}$ Chinese dim sum restaurant	28.7	
	Hong Kong	$\text{PM}_{2.5}$ Western Canteen	21.9	
Levy et al. 2002	USA	$\text{PM}_{2.5}$ food court	200	1.4×10^5
Wallace et al., 2004	USA	Cooking dinner		1.3×10^4
		Cooking breakfast		5.7×10^3

Table 3 Cont. Particle mass and number concentration measured in indoor environments close to cooking activities

Reference	Location	Comment	Concentration ($\mu\text{g}/\text{m}^3$)	Particle Number concentration (part/cm^3)
He et al., 2004a	Australia	PM _{2.5} (48h) cooking	37	1.27×10^5
		PM _{2.5} (48h) cooking pizza	735	1.37×10^5
		PM _{2.5} (48h) frying	745	1.54×10^5
		PM _{2.5} (48h) grilling	718	1.61×10^5
		PM _{2.5} (48h) kettle	13	1.56×10^4
		PM _{2.5} (48h) microwave	16	1.63×10^4
		PM _{2.5} (48h) oven	24	6.15×10^4
		PM _{2.5} (48h) stove	57	1.79×10^5
		PM _{2.5} (48h) toasting	35	1.14×10^5
		PM _{2.5} residential kitchen	535.4	2.86×10^4
He et al., 2004c	China	PM _{2.5} Hunan restaurant	1406	
	China	PM _{2.5} Cantonese restaurant	672	
See and Balasubramanian, 2006a, See and Balasubramanian, 2008	Singapore	PM _{2.5} Steaming	66 ± 7.6	5.4×10^4
		PM _{2.5} Boiling	81 ± 9.3	6.9×10^4
		PM _{2.5} Stir-Frying	120 ± 13	9.3×10^4
		PM _{2.5} Pan-Frying	130 ± 15	11×10^4
		PM _{2.5} Deep-Frying	190 ± 20	59×10^4
See and Balasubramanian, 2006b	Singapore	Stir-fry in a wok typical Chinese food commercial food stall PM _{2.5}	286	7.7×10^5
See et al., 2006	Singapore	PM _{2.5} Chinese stall	202 ± 141	
		PM _{2.5} Malay stall	245 ± 77	
		PM _{2.5} Indian stall	187 ± 44	
		PM _{2.5} Background	29 ± 8	
Hussein et al., 2006	Czech Republic	Cooking in a stove, frying, oven		$0.6-1.8 \times 10^5$
Sjaastad et al., 2008	Norway	Frying Beefsteak		1.2×10^3 (a)
Yeung and To, 2008	Hong Kong	Frying vermicelli with beef		89×10^5
		Pan-frying steaks		8.5×10^5
		Pan-frying chicken fillets		8.5×10^5
		Pan -frying pork chops		8.8×10^5
		Hot oil test		6.4×10^5

Table 3 Cont. Particle mass and number concentration measured in indoor environments close to cooking activities

Reference	Location	Comment	Concentration ($\mu\text{g}/\text{m}^3$)	Particle Number concentration (part/cm^3)
Buonanno et al., 2009	Italy	Grilling in a gas stove at maximum power		
		Cheese	283	1.1×10^5
		Wurstel sausage	352	1.3×10^5
		Bacon	389	1.0×10^5
		Eggplant	78	1.2×10^5
		Frying 50 g of chips in a gas stove at maximum power with	118	1.2×10^5
		Olive oil		
		Peanut Oil	68	1.2×10^5
		Sunflower Oil	60	1.1×10^5
		Frying 50 g of chips using an electrical pan with		
sunflower oil	12	1.4×10^4		
olive Oil	27	2.6×10^4		
peanut Oil	13	1.5×10^4		
Buonanno et al., 2010	Italy	PM ₁ range	10-327	$1.1-9.8 \times 10^5$
		PM _{2.5}	12-368	
		PM ₁₀	15-482	
Buonanno et al., 2011	Italy	Grilling 100 g cheese		1.8×10^5
		Frying 100 g cheese		2.8×10^5
		Grilling 100 g bacon		2.0×10^5
		Frying 100 g bacon		2.8×10^5
		Grilling 100 g pork meat		1.6×10^5
		Frying 100 g pork meat		2.3×10^5
		Grilling 100 g eggplant		1.6×10^5
		Frying 100 g eggplant		2.3×10^5
		Grilling 100 g chips		1.5×10^5
		Frying 100 g chips		2.3×10^5
		Grilling 100 g onion		1.6×10^5
		Frying 100 g onion		2.4×10^5
		Glytsos et al. 2010	Czech Republic	Frying a slice of onion with olive oil – electric griddle

Table 3 Cont. Particle mass and number concentration measured in indoor environments close to cooking activities

Reference	Location	Comment	Concentration ($\mu\text{g}/\text{m}^3$)	Particle Number concentration (part/cm^3)
Huboyo et al., 2011	Japan	Tofu boiling	22.8 (1.21-294)	$6.8 \times 10^{2(a)}$
		Tofu frying	41.2 (1.76-707)	$3.0 \times 10^{2(a)}$
		Chicken boiling	30.8 (5.36-1,082)	$2.5 \times 10^{2(a)}$
		Chicken frying	101.6 (1.67-1,366)	$1.1 \times 10^{2(a)}$
To and Yeung, 2011	Hong Kong	Frying vermicelli with beef – gas cooking (Domestic kitchen) – PM_{10}	1,330	
		Frying vermicelli with beef – electric cooking (Domestic kitchen) – PM_{10}	1,030	
		Pan Frying of meat – gas cooking (Domestic kitchen) – PM_{10}	1,020	
		Pan Frying of meat – electric cooking (Domestic kitchen) – PM_{10}	520	
		Deep frying of chicken wings – gas cooking (Domestic kitchen) – PM_{10}	890	
		Deep frying of chicken wings – electric cooking (Domestic kitchen) – PM_{10}	680	
		Deep frying of tofu – gas cooking (Commercial kitchen) – PM_{10}	4,720	
		Deep frying of tofu – electric cooking (Commercial kitchen) – PM_{10}	3,980	
		Griddle frying of meat – gas cooking (Commercial kitchen) – PM_{10}	2,260	
		Griddle frying of meat – electric cooking (Commercial kitchen) – PM_{10}	2,600	

(a) Particles with diameter $0.3 \mu\text{m} < D_p < 0.5 \mu\text{m}$

Table 4. Size distribution studies for cooking aerosols

Study and Country	Location and duration	Sampling method (a)	Food	Environmental condition
Hildemann et al., 1991a USA	Commercial scale kitchen Sampling port located above the cooking surface, below the extractor fan.	Electrical Aerosol Analyser TSI 3030	Meat cooking during frying and charbroiling extra-lean and regular hamburger meat	Mechanical ventilation
Li et al., 1993 Taiwan	Domestic kitchen with a gas stove Sampling ports 3m away from the gas stove	DMA TSI 3932; CPC TSI 3022	scrambling eggs, frying chicken, and cooking soup	Windows and doors were closed during measurements
Siegmann and Sattler, 1996 Switzerland	Laboratory kitchen Hot oil at 223, 236 and 256°C.	SMPS	Rapeseed oil	Closed window
Abt et al., 2000 USA	Domestic kitchen with gas and electric stoves. Samples collected over 6-day periods Equipment located in an indoor location adjacent to the kitchen.	SMPS TSI 3934; Electrostatic classifier TSI 3071A; CPC TSI 3022a; APS TSI 3310A	Frying, sautéing, barbequing, oven cooking and toasting	Open doors
Dennekamp et al., 2001 UK	Laboratory kitchen with gas and electric stoves Sampling inlet at face level in front of the cooker	SMPS TSI 3934; CPC TSI 3022A	Vegetable oil used to stir-fry 500 g of vegetables and also 5 rashers of bacon	No ventilation. All windows and doors were closed.
Wallace et al., 2004 USA	Domestic kitchen using gas stove Measurements performed in the duct of the ventilation system.	DMA Electrostatic classifier TSI 3071; CPC TSI 3010; APS TSI 3320; Optical particle counter model 500-I Climet Instruments	Deep frying (peanut oil) of flour tortillas; stir fry (peanut oil) vegetables and frying eggs with butter.	No ventilation. Forced ventilation (recirculation of air)

(a)DMA - Differential Mobility Analyser; SMPS - Scanning mobility particle sizer; APS - Aerodynamic particle sizer; CPC - Condenser Particle Counter

Table 4 Cont. Size distribution studies for cooking aerosols

Study and Country	Location and duration	Sampling method (a)	Food	Environmental condition
Wallace et al., 2006 USA	Personal and indoor (living room) measurements for 7 days in free-style living conditions.	Real time concentrations: Personal and indoor sampling using optical particle counter (personal MIE DataRAM) Integrated exposure: Personal – PEM gravimetric monitor Indoor – Harvard impactor monitor	Normal cooking activities	No control on ventilation
Hussein et al., 2006 Czech Republic	Domestic kitchen using an electrical stove and adjacent living room. Continuous measurement for 15 days at 3 min intervals Sampling ports at 1.5m from the ground and 1m (kitchen) and 5m (adjacent room) from the stove.	SMPS TSI 3934C;	Normal cooking activities (e.g. boiling potatoes, soup, rice, pasta, frying potatoes or pancakes, toasting and baking chicken in the oven.	Natural ventilation
See and Balasubramanian, 2006a, b Singapore	Domestic kitchen Inlets located 0.5 m above the gas stove	SMPS TSI 3034	Steaming, boiling, pan-frying, stir-frying, and deep-frying a pack of 150 g plain tofu (soybean curd) using corn oil.	No ventilation. All windows and doors were closed.
Yeung and To, 2008 China	Commercial kitchen with 2 gas stoves, and an electric griddle	SMPS TSI 3934; electrostatic classifier TSI 3071A; CPC TSI 3022,	Stir frying Chinese food and frying western food	No ventilation. All windows and doors were closed.
Sjaastad et al., 2008 Norway	Laboratory kitchen (19m ²) with electric stove in the middle of the floor with kitchen hood and adjoining room. Sampling ports 1m above the floor (all location) and 1.3 away from the stove (in the kitchen).	Kitchen: Particle counter Met One Model 237B; SMPS TSI-3936 Adjoining room : Electrostatic classifier TSI-3080 Ultrafine CPC TSI-2025A	Frying a beef steak with margarine at maximum power.	Mechanical ventilation.

(a) DMA - Differential Mobility Analyser; SMPS - Scanning mobility particle sizer; APS - Aerodynamic particle sizer; CPC - Condenser Particle Counter

Table 4 Cont. Size distribution studies for cooking aerosols

Study and Country	Location and duration	Sampling method (a)	Food	Environmental condition
Yeung and To, 2008 Hong Kong	Laboratory kitchen (168m ³) with gas stove and electric griddle. Fume hood installed above cooking area.	SMPS TSI 3734; Electrostatic classifier TSI 3071A; CPC TSI 3022A	Chinese style – frying vermicelli with beef- in gas stove; Western style – pan-frying steaks, chicken fillets or pork chops - in electric griddle and hot oil in electric griddle.	No ventilation
Buonanno et al., 2009 Buonanno et al., 2011 Italy	Open plan laboratory kitchen (80m ²) using gas and electrical stoves. Sampling 2 meters away from the stove for 8-10 mins.	SMPS TSI 3936 APS TSI 3321 CPC TSI 3775 Nanometer Aerosol sampler (TSI 3089)	Fry and grill different ingredients: pork meat, eggplant, chips and cheese, bacon and oils (olive oil, peanut oil and sunflower oil)	Minimum ventilation - doors and windows closed. Normal ventilation -doors and windows closed with mechanical ventilation in operation.
Buonanno et al., 2010 Italy	15 pizzerias Sampling 2 meters away from the stove for 8-10 mins.	SMPS TSI 3936 APS TSI 3321 CPC TSI 3775 Nanoparticle surface area monitor TSI 3550 ; PM ₁₀ , PM _{2.5} and PM ₁ measured using a DustTrak DRX Aerosol Monitor TSI 8534	Baking pizza	Normal commercial kitchen ventilation.
Glytsos et al., 2010 Czech Republic	Laboratory room (60m ³) Electric stove Sampling ports 0.9 m above the floor.	DustTrak Aerosol Monitor TSI 8520; P-Trak Ultrafine particle counter TSI 8525 ; GRIMM SMPS+C system - GRIMM, CPC Model 5.403 and Long Vienna DMA.	Frying half of an onion diced in hot olive oil (15 mL).	Mechanical ventilation using the air conditioning system.
Huboyo et al., 2011 Japan	In a kitchen (8.5 m ²) with fumehood and adjoining room (3 m ²). Cooking with single gas stove at medium setting. Sampling ports 1.1m away from the stove (in the kitchen) and 5 m away in the adjoining room.	Sioutas cascade Impactor (SKC); PM _{2.5} UCB optical particle counter (Barkeley Air Monitoring Group)	Frying in sunflower oil and boiling 400 g of soybean curd (tofu) and 400 g of chicken	Ventilation system (standard exhaust fan); Natural ventilation (windows opened)

(a) DMA - Differential Mobility Analyser; SMPS - Scanning mobility particle sizer; APS - Aerodynamic particle sizer; CPC - Condenser Particle Counter

Table 5. Particle diameter mode (i.e. diameter representing highest particle number concentration) of particle number size fraction distribution from cooking activities

Reference	Location	Comment	Diameter (nm)
Li et al., 1993	Taiwan	Frying Chicken	30-50
Siegmann and Sattler, 1996	Switzerland	Rapeseed Oil	30-100
Kleeman et al., 1999	USA	Meat charbroiling	180-320
Abt et al., 2000	US	Size range Increasing diameter during cooking Oven cooking event	20-70
Wallace et al., 2004	USA	Cooking dinner	18-50
		Cooking breakfast	10-50
Yeung and To, 2008	Hong Kong	Frying vermicelli with beef	140
		Pan-frying steaks	150
		Pan-frying chicken fillets	115
		Pan –frying pork chops	102
		Hot oil test	107
Buonanno et al., 2009	Italy	Grilling in a gas stove at maximum power	
		Bacon	50
		Cheese	40
		Eggplant	20
		Wurstel sausage	40
		50 g of chips fried with sunflower oil	50
		50 g of chips fried with olive Oil	61
		50 g of chips fried with peanut Oil	50
50 g bacon grilled on a gas stove	60		
Glytsos et al., 2010	Czech Republic	Frying a slice of onion with olive oil	20 – 45
Buonanno et al., 2011	Italy	Frying 100 g mozzarella	80
		Frying 100 g chips	60
		Grilling 100 g bacon	90
		Grilling 100 g eggplant	40

Table 6. Sampling, extraction and analysis of emissions from cooking

STUDY & RESEARCH OBJECTIVES	SAMPLING CONDITIONS	SAMPLE SUBSTRATE PRE-TREATMENT	EXTRACTION PROCEDURE	ANALYTICAL METHODOLOGY	COMPOUNDS ANALYSED
Rogge et al., 1991 Characterise organic compound composition emitted during meat charbroiling	1.8 µm cyclone upstream of 3 pumps Flow rate 9.0-9.6 L/min Sampling duration: 70-80 min	47mm teflon and quartz fiber	Samples were a composite of 15 quartz filters Extraction: Hexane (two times) and with benzene:2- propanol (2:1; three times) Extraction method: mild sonication Final volume reduced to 200-500 µL . Derivatization: one aliquot of the extract was derivatized with diazomethane to convert organic acids to their methyl ester analogues	GC/MS 30-m column	N-alkanes, branched alkanes, alkenes, alkynes, ketones, carbonyls, aromatic hydrocarbons, lactones, amides, saturated and unsaturated fatty acids, dicarboxylic acids, furans amides, steroids.
Wu et al., 1998 Determination of mutagenic PAH emitted from cooking oils	Personal sampling pump Flow rate: 2 l/min Sampling duration: 30 min.	37-mm Grade AA glass fiber filter paper	Extraction with a 200 ml acetone then concentrated to 10 ml in a vacuum rotary evaporator and evaporated to dryness under nitrogen stream. Residue was redissolved in 2ml for analysis.	HPLC system (LH-20 column 15 mm id=190 mm) for PAHS. For detection of aminopyrenes a HPLC Hewlett Packard 1050 was used equipped with a 25-cm Nucleosil C column and spectrofluorimeter.	polycyclic aromatic hydrocarbons; nitro-polycyclic aromatic hydrocarbons
Schauer et al., 1999a Characterise organic compound composition emitted during meat charbroiling Schauer et al., 2002 Characterise organic compound composition emitted during oil cooking	Emissions sampled in the ventilation system of a commercial kitchen downstream of the filter and grease extractor. Sampling time was 85 min. Dilution tunnel: mix exhaust emissions with 25- to 180-fold excess of HEPA filtered air. 1.8 µm AIHL-design cyclone separators upstream of samplers. Flow rate in each sampling train was 10 L/min, except sampling train a) at 30 L/min and sampling train g) at 0.2 L/min. Organic compounds collected using: a) 1 XAD coated denuder upstream of 3 quartz filters in parallel followed by 2 PUFs in series. b) 3 quartz filters followed each by 2 PUFs in series. EC/OC collected using: c) 2 quartz filters in series Mass emissions, trace metals and organic acids collected using: d) Teflon filter upstream of two KOH impregnated quartz fibre filters Mass emissions & soluble ions collected using: e) Teflon filter VOC collected using: f) 6-L SUMA canister downstream of teflon filter e) Carbonyls collected using: g) DNPH-impregnated C18 cartridges	Quartz fibre filters prebaked at 550°C for 12 h Denuders coated following protocol described in Gundel et al. (1995) (Gundel et al., 1995) PUF plugs were pre-cleaned with 4 successive extractions of Dichloromethane/acetone/hexane (2:3:5). Solvent was removed by compressing the PUFs. Plugs were air dried in a dark organic free room, and stored in pre-cleaned glass jars at -20°C.	<i>Quartz fibre filters:</i> Extraction: Hexane (two times) followed by benzene/2- propanol (2:1; three times) Extraction method: mild sonication <i>Denuders and PUFs:</i> Extraction: Dichloromethane/acetone/hexane (2:3:5) (4 times) Extraction method: Manual shaking In all cases, extracts from each aliquot were combined and concentrated to 250 µL. Concentrated extract was split in two. Derivatisation: one aliquot of the extract was derivatised with diazomethane to convert organic acids to their methyl ester analogues. C18 cartridges were extracted as described in Grosjean et al. (1996) (Grosjean et al., 1996) Teflon filters were extracted in water for water-soluble ions.	Organic compounds: Denuder, filter and PUF extracts (derivatized and undervatized aliquots) were analysed by GC/MS Hewlett Packard 5890 series fitted with a 30m, 0.25 mm inner diameter, 0.25 µm film thickness HP-1701 capillary column. Total non-methane organic gases and individual VOCs (C1-C10) were analysed from the SUMA canisters by GC/FID as described in Fraser et al. (1997) (Fraser et al., 1997) Carbonyl collected in the C18 cartridges were eluted with 2 mL acetonitrile analysed by LC/UV as described by Grosjean et al. (1996). Organic and elemental carbon (EC/OC) as described by Birch and Cary (1996) (Birch and Cary, 1996) Trace metals were analysed by XRF. Soluble ions were analysed by AA and IC.	N-alkanes, branched alkanes, alkenes, alkynes, ketones, carbonyls, aromatic hydrocarbons, lactones, amides, saturated and unsaturated fatty acids, dicarboxylic acids.

Table 6 Cont. Sampling, extraction and analysis of organic emissions from cooking

STUDY & RESEARCH OBJECTIVES	SAMPLING CONDITIONS	SAMPLE SUBSTRATE PRE-TREATMENT	EXTRACTION PROCEDURE	ANALYTICAL METHODOLOGY	COMPOUNDS ANALYSED
<p>Svendsen et al., 2002 Characterise aldehydes and fat aerosol collected in the breathing zone of the cook in fumes from commercial restaurants.</p>	<p>Personal exposure sampler with inlets located in the shoulder of the cook of 19 commercial kitchens using deep frying devices equipped with ventilation hoods.</p> <p>Aldehydes were collected a sampling device containing silica impregnated with 2,4-dinitrophenyl hydrazine. Flow rate was 1.5 L/min during 1.5-2.5 hours.</p> <p>Fat aerosol collected onto pre-weighted one glass fibre filter (Nucleopore AAA). Flow rate, 2 L/min during 65 to 200 mL.</p> <p>Total number concentration was measured with TSI 3936 SMPS used to measure the</p> <p>PAHs were collected onto glass fibre filters in a filter holder and 2 XAD-2 tubes downstream. Flow rate, 1 L/min during 200 min.</p>	<p>None</p>	<p>Fat aerosol extracted with 5 mL of 1,1,2-trichloro-1,2,2-trifluoroethane.</p> <p>The aldehydes were reacted with 2,4-dinitrophenylhydrazine (DNPH) to form the corresponding stable hydrazone derivatives. The derivatives were eluted with HPLC grade acetonitrile.</p>	<p>Fat aerosol was determined using a FT-IR (Perkin Elmer 1605).</p> <p>The eluate was injected onto a C18 reverse phase column and detected using a UV detector operating at 360 nm.</p>	<p>Aldehydes, fat aerosol</p>
<p>McDonald et al., 2003 Characterise organic compound emission composition emitted during charbroiling and grilling of chicken and beef</p>	<p>University lab kitchen following commercial standard procedures. Emissions collected at the end of a residence chamber to allow the gas/particle equilibrium.</p> <p>2.5 µm cyclone separators upstream of samplers.</p> <p>Flow rate in each sampling train was 113 L/min.</p> <p>Samples collected on Teflon filter for PM_{2.5} and elements.</p> <p>Samples collected on quartz filters for carbon and ion analysis</p> <p>Samples collected on Teflon-impregnated glass fibre (TIGF) filter followed by a PUF/XAD-4/PUF sandwich cartridge for speciated particle-phase and semi-VOCs.</p> <p>CO was measured using non-dispersive infrared analyser.</p>	<p>Quartz fibre filters were baked at 900°C for several hours.</p> <p>XAD-4 was solvent extracted in a Soxhlet with methanol followed by dichloromethane.</p> <p>TIGF filters were cleaned by sonication in CH₂Cl₂ for 30 min followed by another 30 min sonication in methanol.</p> <p>PUFs were rinsed with distilled water and Soxhlet extracted with hexane/ether (90:10), followed by acetone.</p>	<p>Half of the quartz fibre filter was extracted with 10 mL of distilled-deionised water.</p> <p>TIGD filters and PUF/XAD-4/PUF sorbent were solvent extracted and combined for analysis.</p>	<p>PM_{2.5} mass determined by gravimetric analysis.</p> <p>Ionic species determined by ionic chromatography. NH₄⁺ was analysed by indophenol automated colorimetry. Water-soluble K⁺ was analysed by atomic absorption spectrometry.</p> <p>Carbon by thermal/optical reflectance. 0.56 cm² punch was analysed for OC/EC by the TOR method.</p> <p>Elements by X-ray fluorescence.</p> <p>Organics determined with an Agilent GCMS (GC Model 6890plus, MSD Model 5973N) equipped with a 60m x 0.25mm x 0.25 µm DB5-MS capillary column.</p>	<p>PM_{2.5}, CO, OC/EC, inorganic species, elements, lactones, sterols, PAHs, biphenyls,</p>
<p>Zhu and Wang, 2003 Characterise PAH emitted in commercial and domestic Chinese kitchens</p>	<p>A sampler was located in a new kitchen 0.5 m from the pan (cooking methods) and in the centre of the kitchen (domestic and commercial kitchens). In all cases, the sampler was 1.5 m above the ground level. All doors and windows were closed during cooking. Electric hobs were used for cooking.</p> <p>Samples were collected over 100 mins to test different cooking methods, and over 2 consecutive days for 12-h (8:00 – 20:00) in domestic and commercial kitchens.</p> <p>Low noise small samplers (MP-15CF) operated at 1.0 l/min equipped with a Whatman glass filter (GFF, 25 mm) collected particle bound PAHs and a XAD-2.5 g cartridge collected the gaseous PAHs.</p>	<p>Filters were combusted overnight and sealed in aluminium foil.</p> <p>XAD-2 cartridges were pre-extracted in dichloromethane and methanol for 48 h, vacuum-dried in desiccators and stored in solvent rinsed glass jars.</p>	<p>Extraction by sonication for 30-min with a 20 ml mixture of DCM and acetonitrile (3:2). The extract was concentrated to 10 ml and 30 µl of dimethyl sulfoxide was added. The mixture was concentrated under nitrogen and 1ml of methanol was added. 100 µl were injected for analysis.</p>	<p>HPLC (Hitachi L-7000 series) consisting of a precolumn (Supelco, 5C-18, 4.6x 50 mm) for PAH condensation and cleanup, a main column (Wakosil, 5C-18, U4:6 250 mm) for separation and a fluorescence detector.</p>	<p>PAHs</p>

Table 6 Cont. Sampling, extraction and analysis of organic emissions from cooking

STUDY & RESEARCH OBJECTIVES	SAMPLING CONDITIONS	SAMPLE SUBSTRATE PRE-TREATMENT	EXTRACTION PROCEDURE	ANALYTICAL METHODOLOGY	COMPOUNDS ANALYSED
Chen and Chen, 2003 Characterise PAHs in fumes during frying of chicken.	Emissions collected on adsorption wool fitted on the cover of frying tank (closely tight during sampling)	Adsorption wool	Soxhlet extraction for 20hrs using acetone to 1ml then evaporated to dryness then residue dissolved in 10 ml acetone and stored for GCMS analysis.	GC/MS equipped with an HP-5MS (30 m x 0.25 mm i.d., 0.25 um film thickness)	PAHs
Li et al., 2003 Characterise PAHs in fumes during cooking of different styles	Emissions collected isokinetically from the exhaust vent in commercial kitchens. Three consecutive samples were collected at 10L/min for 45 min during the cooking time. Particle bound PAHs were collected on a tube-type glass fibre thimble (25x90 mm). Gaseous PAHs were collected onto a 5-cm polyurethane foam (PUF) followed by a 2.5 cm Xad-16 resin cartridge supported by a 2.5 cm PUF.	Samples were kept prior and after sampling in cleaned screw-capped glass bottles and jars.	Samples were extracted in a Soxhlet extractor with 1L of mixed solvent n-hexane/dichloromethane (1:1) for 24 hours. The extract was concentrated, cleaned and re-concentrated to 1 or 1.5 mL.	Hewlett-Packard GC HP 5890A with a Mass Spectrometer detector HP 59H72 equipped with a HP Ultra 2 50m x 0.32 mm x 0.17 um column.	PAHs
He et al., 2004b Characterise fumes emitted during Chinese style cooking	Samples collected at the exit of the exhaust vent of two commercial kitchens. Sampling times were 90-120min at lunchtime and dinner. Samples collected onto two honeycomb sampler and a three stage cascade impactor to collect PM _{2.5} at 25 L/min. One honeycomb contained PTFE filters for particle mass determination and and ionic species analysis. The second honeycomb and the cascade impactor were loaded with quartz filters (Pallflex 2500QAT-UP) for the determination of EC/OC and organic speciation.	Quartz fiber filters were baked for 4 hours at 500°C. Pre- and post-sampling filters were stored in pre-cleaned 250 mL glass jars with 3-5 mL of methylene chloride to prevent microbial growth. Sampled filters stored in the freezer.	Samples extracted with dichloromethane (3 times) and methanol (2 times) for 20 min using a mild ultrasound bath. Reduced to 5 mL with rotary evaporation and further concentrated to 1ml under a N ₂ stream and split into three fractions. Two fractions were derivatised with BF ₃ /CH ₃ OH and bis-(trimethylsilyl) trifluoroacetamide (BSTFA) plus 1% trimethylchlorosilane (TMCS) to convert organic acids and unmethylated compounds to their methyl ester and trimethylsilyl derivatives respectively. Derivatisation temperatures and times were 80°C for 30 min and 85°C for 40 min respectively.	PM _{2.5} mass determined by gravimetric analysis. Ionic species determined by ionic chromatography (DX-600, Dionex Corp). EC/OC determined with the Sunset analyser. Organics determined with an Agilent GCMS (GC Model 6890plus, MSD Model 5973N) equipped with a 60m x 0.25mm x 0.25 um DB5-MS capillary column.	N-alkanes, n-fatty acids and dicarboxylic acids; PAHs and other compounds including cholesterol and levoglucosan.
He et al., 2004c Characterise fumes emitted during Chinese style cooking	Samples collected at 40-60 cm at leeway from the exhaust vent of two commercial kitchens. Sampling times were 100-120min at lunchtime, and 45 minutes at dinner. Samples collected onto quartz fibre filters with a three stage cascade impactor (<10um, 10-2.5 um and <2.5 um) at 25 L/min.	Quartz fiber filters were baked for 2 hours at 500°C. Pre- and post-sampling filters were stored in pre-cleaned 250 mL glass jars with 3-5 mL of methylene chloride to prevent microbial growth. Sampled filters were stored in the freezer.	Samples extracted with methylene chloride (3 times) for 20 min using a mild ultrasound bath. Reduced to 5 mL with rotary evaporation and further concentrated to 1ml under a N ₂ stream.	GC/MS Autosystem XL Gas Chromatography coupled with a TurboMass Mass spectrometry (Perkin Elmer) equipped with a 60m x 0.32mm x 0.25 um fused silica capillary column (PE-35MS)	N-alkanes, n-alkenes, n-fatty acids; n-alkanal; n-alkenals; PAHs

Table 6 Cont. Sampling, extraction and analysis of organic emissions from cooking

STUDY & RESEARCH OBJECTIVES	SAMPLING CONDITIONS	SAMPLE SUBSTRATE PRE-TREATMENT	EXTRACTION PROCEDURE	ANALYTICAL METHODOLOGY	COMPOUNDS ANALYSED
<p>See et al., 2006; See and Balasubramanian, 2006b Characterise PAH and metal composition emitted during Chinese, Malay and Indian style commercial cooking</p> <p>See and Balasubramanian, 2006a, 2008 Characterise emissions from 5 types of cooking methods (steaming, boiling, stir-frying, pan-frying and deep-frying)</p>	<p>Sample collected at 1.5m above ground level at the opposite site of a 4 LPG burners stove in commercial food stalls. (See et al., 2006; See and Balasubramanian, 2006b)</p> <p>Sample collected at 1.5m above ground level and 0.2 m from a 2-burner domestic stove with no ventilation. Samples collected during cooking activities. (See and Balasubramanian, 2006a, 2008).</p> <p>Samples collected for 12 hours during cooking and non-cooking activities.</p> <p>A MiniVol portable air sampler (Airmetrics) collected PM_{2.5} at a flow rate of 5 L/min onto:</p> <ul style="list-style-type: none"> - 47mm 2 µm PTFE Teflon filter for gravimetric, metals and ion analysis. - 47mm QMA quartz filters for PAH 	<p>QMA filter was pre-combusted at 400°C for 24h prior to sampling.</p> <p>No pre-treatment of Teflon filter</p>	<p><i>PAH</i> Microwave extraction using 20mL acetone:hexane (1:1) for 20 min at 150W. Extracts concentrated to 3 mL using a rotary evaporator. Further concentration to almost dryness with N₂ stream and reconstituted with 1 mL of extraction solvent.</p> <p><i>Metals</i> Microwave extraction as described by Swami et al. (2001) (Swami et al., 2001)</p> <p><i>EC/OC</i> 2 6mm punches of a quartz fibre filter. One punch was combusted at 350°C for 24h to remove the OC.</p>	<p><i>PAHs</i> Hewlett Packard 6890 series GC/MS fitted with a DB-5MS 5%-phenyl-methylpolysiloxane 30m x 0.2 mm internal diameter x 0.25 µm film thickness.</p> <p><i>Metals</i> Perkin Elmer ELAN 6100 ICP/MS</p> <p><i>EC/OC:</i> Both combusted and uncombusted punches were analysed for carbon using a 2400 Series II CHNS/O analyser (Perkin Elmer) operated at the CHN mode with acetanilide as calibration standard.</p>	<p>PAHs</p> <p>Metals</p> <p>EC/OC</p>
<p>Zhao et al., 2007a, b Characterise organic compound emission composition emitted during Chinese and Western style cooking</p>	<p>Emissions sampled at the exhaust vent of the ventilation system of commercial kitchens downstream of the filter treatment methods. Samples collected during rush hour at lunch and dinner times. Sampling time was 120 min.</p> <p>2 medium-volume samplers at a flow rate of 78 L/min collected samples in 90mm quartz fibre filter.</p> <p>2 Dustraks (TSI) monitored the relative concentrations of PM_{2.5} and PM₁₀. Background PM_{2.5} was collected in the city using a hi-volume sampler (Andersen).</p>	<p>Quartz fiber filters were baked at 450°C for 4.5 hours. Prior to sampling, filters were stored in a freezer.</p>	<p>Extraction with three successive portions of dichloromethane and methanol (3:1) for three 15-min in the ultrasound bath at room temperature. Extracts were filtered and distilled under negative pressure to 3-5 mL, subsequently concentrated to 1 mL under N₂ gas stream, and divided in three portions: Portion 1 – analysed directly in GCMS for non polar organic compounds. Portion 2 - Derivatized with BSTFA plus 1% TMCS at 70°C for 2 h. This was analysed for polar organic compounds. Third portion – stored at 4°C as a backup.</p>	<p>Organics Agilent 6890plus / MSD model 5973N GC/MS using a DB-5MS 60m x 0.25 mm internal diameter x 0.25 µm film thickness column</p> <p><i>EC/OC:</i> Carbon analyser Sunset Lab.</p>	<p>N-alkanes, PAHs, N-alkanals, N-alkanones, lactones, amides, saturated and unsaturated fatty acids, dicarboxylic acids, anhydrides, sterols</p>
<p>Sjaastad and Svendsen, 2008; Sjaastad et al., 2010; Sjaastad and Svendsen, 2009 Characterise PAHs, aldehydes and particulate matter collected in the breathing zone of the cook in fumes from frying a beefsteak.</p>	<p>Model kitchen (19 m²) containing gas or electric hobs and a canopy fume hood.</p> <p>Personal exposure sampler with inlets located in the shoulder of the cook.</p> <p>PAHs were collected onto glass fibre filters in a filter holder and 2 XAD-2 tubes downstream. Flow rate, 1 L/min during 200 min.</p> <p>Aldehydes were collected into stainless steel sorbent tubes filled with 220 mg Tenax TA. Flow rate, 100 mL/min for 10-200 min.</p> <p>Total particles collected onto pre-weighted double Gelman glass fibre filters. Flow rate, 2 L/min during 65 to 200 mL.</p> <p>Total number concentration was measured with TSI 3936 SMPS.</p>	<p>None</p>	<p>PAH were desorbed in dichloromethane.</p>	<p>PAH measured by a commercial laboratory following a modified version of AMI L5, NIOSH 5515, ISO/CD 12884 and VDI 3873.</p> <p>Aldehydes measured using an automatic thermic desorption unit ATD 400 (Perkin Elmer) connected to a GCMS (Focus GC-DSQ, Thermo Electron Corporation).</p>	<p>PAH; aldehydes</p>

Table 7. Main identified cooking marker species in the literature

COMPOUND ANALYSED	SOURCE IN FOOD	OTHER SOURCES IN THE ENVIRONMENT
Unsaturated fatty acids Oleic acid- 9-octadecenoic acid-meat tracer, canola oil (Schauer et al, 2002) Linoleic acid- 9,12-octadecadienoic acid Palmitoleic acid- 9-hexadecenoic acid meat cooking (Zhao, 2007a; Robinson et al, 2006)	Combustion of triglycerides and phospholipids from seed oil, vegetable oil, fats of animals and meat cooking (Robinson et al, 2006)	Biomass smoke, motor vehicle exhaust and road dust (Robinson et al., 2006)
Saturated Fatty Acids hexanoic acid octanoic acid nonanoic acid-from seed oil (Schauer et al, 2002) hexadecanoic acid, palmitic acid, (Robinson et al, 2006)	Combustion of triglycerides and phospholipids from seed oil, vegetable oil and fats of animals. The acids are formed directly from the pyrolysis of their glycerol ester precursor analogues (nonanoic acid formed from the breakdown of oleic acid present in seed oil (Schauer et al, 2002)	Palmitic acid are emitted also from biomass smoke, motor vehicle exhaust, road tire dust (Robinson et al, 2006), tyre dust cigarette smoke, roofing asphalts and fuel combustion (Nolte et al., 1999)
Dicarboxylic Acids-C4-C8 hexanedioic acid – from meat cooking and seed oil octanedioic acid – from seed oil nonanedioic acid, tetradecanoic acid, octadecadienoic acid from soybeans oil (Schauer et al, 2002)	Products of dialdehydes formed during auto oxidation of unsaturated lipids. Produced from meat cooking (C ₄ -C ₈ high concentrations for hexanoic acid) and heating of seed oil (C ₈ higher concentrations) (Zhao, 2007a).	
Polycyclic Aromatic Hydrocarbons pyrene chrysene –seed oil and meat cooking (Zhao, 2007a) benzo[a]pyrene	Incomplete combustion of organic substance (cooking materials such as meat, vegetables, oil)	House heating Cigarette smoking (Kleeman et al., 2008). Heavier PAH (coronene, benzo[ghi]perylene, indeno[1,2,3-cd]pyrene) are emitted from motor vehicles and retene from biomass burning (Brinkman et al., 2009)
Molecular bio markers • Monosaccharide Anhydrides- from breakdown of cellulose during cooking (Zhao, 2007a) Galactosan Mannosan levoglucosan • Sterols <i>B</i> -sitosterol –present in animal and vegetable body tissue (Zhao, 2007a). Cholesterol – from meat cooking (Zhao, 2007a; Robinson et al, 2006) stigmasterol	From the organic compounds of biological origin which have restricted occurrence and molecular stability so can be detected in body tissues. Plant lipid membranes and waxes. For Chinese food the average ratio of levoglucosan/(mannosan+galactosan) is 12 (Zhao, 2007a).	Levoglucosan is released from wood burning. (Kleeman et al., 2008; Zhao et al., 2007a) Cholesterol produced from cigarette, debris of plant and road dust (Zhao et al., 2007a; Robinson et al., 2006; Nolte et al., 1999).
N Alkanes C₂₃-C₃₁ C ₂₃ -C ₃₁ from cooking material/contents (Zhao, 2007a)	From cooking material/contents (Zhao, 2007a)	From motor vehicles (Brinkman et al., 2009)
Lactones C₇-C₁₈ From food cooking (Zhao, 2007a; Schauer et al., 2002). 5-propyldihydro-2(3H) furanone (Schauer et al, 2002) 5-dodecyldihydro-2(3H) furanone (Schauer et al, 2002)	Meat charbroiling and food cooking (Schauer et al, 2002)	
Alkanals and alkanones C₉-C₁₅ from cooking oil Nonanal (Zhao, 2007a) 2-pentadecanone- from soybean oil and seed oil (Schauer et al, 2002) 2-nonanone 2-undecanone 2-pentadecanone	Combustion of triglycerides in oil (Zhao, 2007a). From the decomposition of unsaturated fatty acids (oleic acid) (Schauer et al, 2002)	
Inorganic elements and ions From meat cooking Aluminium (Schauer et al, 1999a) Silicon (Schauer et al, 1999a) Phosphorus (Schauer et al, 1999a) Sulphur (Schauer et al, 2002; Schauer et al, 1999a) Chlorine (Schauer et al, 1999a) Potassium (Schauer et al, 1999a) Sodium (Schauer et al, 2002; Schauer et al, 1999a) Nitrate (Schauer et al, 2002; Schauer et al, 1999a)	From meat, vegetables and sauces (Schauer et al, 2002)	From soil, motor vehicles, cigarettes and biomass burning (Brinkman et al., 2009)

Table 8. Concentrations of organic compounds from western-style fast food and from Chinese cooking (ng/mg of particulate organic matter) (Zhao et al., 2007b,c)

Organic compounds	Western-style fast food cooking	Chinese cooking
n-Alkanes	3860	1880
Polycyclic aromatic hydrocarbons	40	2860
n-Alkanones	22700	2440
n-Alkanals	29200	3440
Lactones	13300	2140
Amides	4690	531
Saturated fatty acids	374700	26800
Unsaturated fatty acids	93300	29030
Dicarboxylic acids	57900	2050
Monosaccharide anhydrides	97	314
Sterols	487	1680

Table 9. Polycyclic aromatic hydrocarbons emitted from cooking food – indoor concentrations and emission factors (see columns for units)

PAH	Rogge et al, 1991		Schauer et al, 1999a	Schauer et al, 2002						Zhu and Wang, 2003								McDonald et al, 2003						
	frying hamburger	charbroiling hamburger	Hamburger meat charbroiled	stir fry in soybean oil		stir fry in canola oil		deep frying of potatoes in hydrogenated oil		Boiling Fish	Frying Fish	Broiling Fish	Boiling short pork chops	Frying short pork chops	Broiling short pork chops	Commercial Kitchen	Domestic Kitchen	Hamburger Auto-Char	Hamburger Under-Char	Steak Under-Char	Chicken Under-Char	Hamburger Griddle	Chicken Griddle	
	mg/kg		µg/kg of cooked meat	µg/kg of cooked vegetables		µg/kg of cooked vegetables		µg/kg of cooked potatoes		µg/m ³ /kg								µg/m ³		mg/kg				
	particle	particle	gas+particle	gas	particle	gas	particle	gas	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	
Naphthalene			8990	645		588		338		0.028	0.25	1.1	0.045	0.53	1.3	2.33	1.84	23.04	19.11	14.8	8.75	0.61	1	
Acenaphthene																		0.28	0.15	0.15	0.1	0.02	0.05	
Acenaphthylene				38		37		19		nd	0.08	0.1	0.0013	0.57	2.4	4.20	1.24	4.89	4.24	4.28	2.06	0.16	0.13	
Fluorene										nd	0.042	0.13	nd	0.071	0.34	0.94	0.07	1.09	1.26	1.17	0.72	0.21	0.18	
Phenanthrene			1220	138	7	120	8	83	2	0.0021	0.045	0.043	0.0031	0.069	0.18	0.36	0.63	4.88	4.88	5.31	3.46	2.07	1.87	
Anthracene			160	11	1	8	2	6	2	0.003	0.048	0.043	0.0022	0.033	0.13	0.43	0.28	0.91	0.94	1.03	0.88	0.17	0.44	
Fluoranthene	0.13	0.35	520	40	7	24	5	19	1	0.011	0.25	0.15	0.015	0.31	0.43	1.08	0.94	0.88	1.4	1.28	1.28	0.86	0.62	
Pyrene	0.09	0.74	670	28	4	15	5	19	1	0.009	0.25	0.15	0.013	0.17	0.55	4.20	0.63	1.15	1.9	1.56	1.8	1.15	0.82	
Retene																		0.01	0.02	0.01	0.03	0.02	0.02	
Benzo[ghi]fluoranthene										0.0082	0.18	nd	0.012	0.39	1.4	0.39	0.27							
Benzo[a]anthracene	0.02	0.29								0.0022	0.033	0.028	0.0031	0.049	0.14	0.76	0.23	0.22	0.22	0.11	0.34	0.07	0.12	
Chrysene	0.11	0.95	600 (b)	5 (b)	9(b)		13(b)	5(b)	9(b)									0.24(b)	0.26(b)	0.12(b)	0.29(b)	0.06(b)	0.12(b)	
Benzo[b]fluoranthene	nd	0.21								nd	0.082	0.037	0.0011	0.14	0.26	0.90	0.01	0.4(c)	0.34(c)	0.17(c)	0.24(c)	0	0.01(c)	
Benzo[k]fluoranthene	0.004	0.27								0.0021	0.011	0.01	0.0031	0.079	0.08	2.16	0.04							
Benzo[a]fluoranthene																								
Benzo[b]pyrene																								
Benzo[e]pyrene	nd	0.19								nd	0.055	0.04	nd	0.075	0.089	0.28	0.02	0.14	0.1	0.04	0.06	0.01	0	
Benzo[a]pyrene	nd	0.19																0.17	0.15	0.07	0.1	0.02	0.01	
Perylene	nd	0.03																						
Indeno[1,2,3-cd]pyrene																		0.1	0.09	0.05	0.06	0	0	
dibenz(a,h)anthracene																		0.02	0.01	0.01	0.01	0	0	
Benzo[ghi]perylene	nd	0.24																0.16	0.17	0.09	0.09	0	0	
Coronene																		0.08	0.1	0.05	0.05	0.01	0.02	

Table 9 Cont. Polycyclic aromatic hydrocarbons emitted from cooking food – indoor concentrations and emission factors (see columns for units)

PAH	Li et al, 2003				He et al 2004b				See et al, 2006			Zhao et al, 2007c		Zhao et al, 2007b				See and Balasubramanian, 2008					
	Chinese	Western	Fast Food	Japanese	Hunan cooking	Cantonese cooking	Hunan cooking	Cantonese cooking	Chinese	Malay	Indian	Western-stylefastfoodcooking	Chinese cooking (average)	Cantonese style	Sichuan style	Dongbei style	Hunan style	Steaming	Boiling	Stir-frying	Pan-frying	Deep-frying	
Units	µg/m ³				ng/m ³		ng/mg of particles emitted		ng/m ³			ng/mg of POM		ng/mg of POM				ng/m ³					
	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle	particle
Naphthalene	36.1	61.3	46.8	39.5					1.9	2.8	3.9							0.17 ± 0.01	0.19 ± 0.01	0.30 ± 0.03	0.37 ± 0.04	0.50 ± 0.04	
Acenaphthene	1.65	0.89	0.36	0.285					1	3.1	1.1							0.13 ± 0.03	0.42 ± 0.05	0.57 ± 0.07	0.98 ± 0.10	1.17 ± 0.14	
Acenaphthylene	11.7	4.21	1.55	1.25					2.4	5.6	2.7							0.56 ± 0.06	0.75 ± 0.09	1.55 ± 0.13	1.73 ± 0.18	1.80 ± 0.26	
Fluorene	2.32	1.44	1.92	2.38					3.8	9.2	3.9							0.84 ± 0.07	0.89 ± 0.07	1.53 ± 0.14	1.68 ± 0.12	2.62 ± 0.33	
Phenanthrene	6.38	7.15	5.62	7.41			6.2	6.4	11.5	15.7	9.5	2±1	5.5	7±3	6±2	5±1	4±1	3.18 ± 0.23	3.97 ± 0.32	5.11 ± 0.86	5.87 ± 0.87	8.98 ± 0.68	
Anthracene	1.03	1.15	0.225	0.225			0.27	ND	3	6.1	2.6							0.12 ± 0.01	0.16 ± 0.03	0.31 ± 0.03	0.36 ± 0.05	0.43 ± 0.07	
Fluoranthene	1.32	1.35	0.72	0.635	8.8	10.4	6.2	15.5	6.9	30.7	1.6	nd	110.7	12±36	238±97	56±10	28±16	0.86 ± 0.10	1.05 ± 0.14	1.61 ± 0.16	1.64 ± 0.18	1.64 ± 0.13	
Pyrene	1.33	1.59	0.485	0.325	11	18.7	7.8	27.8	10.9	18.1	2.9	14±14	208.3	243±71	466±191	76±19	49±12	0.80 ± 0.10	0.96 ± 0.14	1.30 ± 0.18	1.35 ± 0.13	1.45 ± 0.16	
Retene												nd	6.2	24±14	1±3	nd	nd						
Benzo[ghi]fluoranthene							1.3	6.1				18±10	77.9	160±30	119±49	20±4	13±3						
Benz[a]anthracene	0.814	0.15	0.276	0.015			0.86	2.5	3.8	23.1	1							0.21 ± 0.03	0.30 ± 0.02	0.38 ± 0.07	0.40 ± 0.06	0.44 ± 0.07	
Chrysene	0.502	0.247	0.596	0.104			0.81	2.8	5.8	48.7	1	7±7	26	59±24	23±8	12±3	10±2	0.56 ± 0.04	0.70 ± 0.09	1.05 ± 0.11	1.07 ± 0.11	1.19 ± 0.15	
Benzo[b]fluoranthene	0.873	1.2	0.752	0.403	2.4 (a)	6.5 (a)	1.7(a)	9.7(a)	21.8	122.4	1.9	nd	18.4	48±30	10±3	9±4	7±2	0.29 ± 0.04	0.70 ± 0.06	1.30 ± 0.10	1.30 ± 0.12	1.38 ± 0.13	
Benzo[k]fluoranthene	0.814	1.29	0.528	0.352					3.7	23.1	0.5	nd	14.6	40±35	8±3	6±4	5±3	0.29 ± 0.02	0.37 ± 0.04	0.48 ± 0.04	0.78 ± 0.10	0.94 ± 0.07	
Benzo[a]fluoranthene												nd	2.8	1±17	nd	1±2	nd						
Benzo[b]pyrene												nd	17.5	52±30	7±4	5±6	6±2						
Benzo[e]pyrene	0.977	1.8	0.256	0.351	1	2.8	0.71	4.2															
Benzo[a]pyrene	1.22	1.59	0.296	0.28	0.7	1.4	0.51	2.1	5.6	16	0.9	nd	11.9	39±42	1±2	3±4	4±2	0.20 ± 0.03	0.23 ± 0.02	0.38 ± 0.05	0.49 ± 0.05	0.56 ± 0.05	
Perylene	1.69	1.48	0.216	0.232								nd	0.4	1±4	nd	nd	nd						
Indeno[1,2,3-cd]pyrene	2.81	1.08	0.177	0.736			ND	3.9	24.4	105.9	1.3	nd	19.6	65±58	4±7	6±6	3±6	0.42 ± 0.04	0.87 ± 0.07	1.09 ± 0.14	1.38 ± 0.19	3.47 ± 0.30	
dibenz(a,h)anthracene	1.91	1.35	0.069	ND					2.7	8.3	1.1							0.046 ± 0.008	0.21 ± 0.02	0.52 ± 0.05	0.62 ± 0.06	1.18 ± 0.15	
Benzo[ghi]perylene	1.65	0.648	0.16	0.152			ND	16.7	31.9	170.1	2.1	nd	65.1	196±116	27±11	18±9	19±8	0.36 ± 0.05	0.79 ± 0.06	1.29 ± 0.17	1.62 ± 0.19	4.33 ± 0.47	
Coronene	0.737	0.448	0.504	0.4			ND	16.5				40	2855										

Note: Sjaastad and Svendsen (2009) reported naphthalene concentrations in the breathing area of chefs working at Norwegian à la carte restaurants in the range of 0.05-0.27 µg/m³.

Table 10. Polycyclic aromatic hydrocarbons in fumes from heated oils - indoor concentrations and emission factors (see columns for units) - and in uncooked oil

Reference	Siegmann and Sattler, 1996			Wu et al., 1998			Chiang et al., 1999			Fortmann et al., 2001					Chen and Chen, 2001			Chen and Chen, 2003			Zhu and Wang, 2003		
Cooking conditions	250°C (oil heated in iron pot)			250°C for 30 min (oil heated in iron pot)			250°C ± 10 for 30 min (oil heated in iron pot)			Uncooked					220°C for 2 h (oil heated in reaction tubes with adsorption tubes placed on them)			frying time 1hr			180-270°C for 2h (oil heated in iron pot)		
	PAH	Rape seed	Corn	Peanut	Lard	Soy bean	Peanut	Lard	Soy bean	Peanut	Olive	Peanut	Canola	Corn	Veget able	Soy bean	sunflower	canola	fried chicken legs in			Rape-seed	Soy bean
Units	mg/kg oil			µg/m ³			µg/m ³			ng/g					µg/g of Smoke			µg/m ³ of smoke			ng/m ³		
naphthalene										31.7	13.9	15.5	13.3	17.6	ND	ND	ND	nd	nd	nd	12.08	13.67	15.5
acenaphthylene										ND	ND	ND	ND	ND	ND	21.1 ± 1.4	12.9 ± 1.2	16.06	5.68	6.79			
acenaphthene										19.9	ND	ND	ND	ND	72.5 ± 2.2	14.5 ± 0.2	62.2 ± 3.1	2.08	4.49	3.29	51.26	20.67	73.5
fluorene										1.73	ND	0.21	0.28	0.3	84.9 ± 0.8	21.7 ± 1.7	59.5 ± 2.5	22.71	nd	nd	2.2	2.27	1.55
phenanthrene	14.4	6.92	37.1							10.7	ND	ND	ND	ND	83.2 ± 4.5	33.0 ± 6.6	60.4 ± 4.3	10.44	4.86	19.17	0.22	0.18	0.26
anthracene										1.12	2.6	1.12	1.54	0.56	31.2 ± 2.6	10.3 ± 1.8	61.1 ± 2.6	14.91	3.57	18.13	0.2	0.13	0.44
fluoranthene	4.2	1.14	15.5							4.07	1.28	0.71	0.65	1.64	98.2 ± 6.3	44.0 ± 2.4	78.1 ± 5.4	10.48	7.42	19.86	0.51	0.31	0.67
pyrene	3.77	1.88	0.66							7.1	10.2	1.79	ND	ND	87.5 ± 3.4	40.1 ± 4.6	68.6 ± 3.6	nd	5.43	nd	0.56	0.42	0.66
benzo[a]anthracene	2.4	0.36	8.75	2.3 ± 0.22	2.1 ± 0.5	21.1 ± 0.8	2.2	2.3	1.3	4.49	13.6	6.51	ND	2.22	46.8 ± 1.2	21.5 ± 1.0	39.0 ± 1.0	5.96	3.97	7.11	1.01	1.57	2.25
chrysene	2.6	0.46	8.5							3.29	14.7	ND	ND	2.22	58.0 ± 6.6	24.6 ± 3.1	44.5 ± 5.2	2.98	3.5	nd	0.13	0.2	0.14
benzo[b]fluoranthene	1	0.47	0.32							77.3	72.8	ND	4.68	5.28	29.8 ± 3.1	13.3 ± 1.8	ND	nd	nd	nd			
benzo[k]fluoranthene															30.6 ± 1.1	15.5 ± 1.5	ND	nd	nd	nd	0.49	0.6	0.94
benzo[a]pyrene	7	0.8	0.1	ND	21.1 ± 0.8	19.6 ± 0.5	n.d	19.6	18.3	8.32	24.5	ND	11	4.22	16.1 ± 1.1	5.9 ± 0.9	10.1 ± 0.3	nd	nd	nd	3.06	2.37	3.2
dibenzo[a,h]anthracene				2.0 ± 0.3	2.4 ± 0.4	1.9 ± 0.1	1.9	2.1	1.8	9.26	27.1	ND	0.59	9.2	15.4 ± 1.2	5.8 ± 0.5	8.8 ± 1.3	23.44	nd		0.7	1.22	1.15
benzo[g,h,i]perylene							8.4				5.31	26.6	18.7	3.2	ND	ND	ND						
indeno[1,2,3-c,d]pyrene							9.84				16.2	30.3	2.67	2.03	8.2 ± 2.4	1.6 ± 0.2	9.9 ± 0.8	nd	nd	nd			

Table 11. N-alkanes emitted from cooking food - indoor concentrations and emission factors (see columns for units)

Reference	Schauer et al, 1999a	Schauer et al, 2002						He et al 2004		Zhao et al, 2007b				Zhao et al, 2007c	
Compound class	Hamburger meat charbroiled	stir fry in soybean oil	stir fry in canola oil	deep frying of potatoes in hydrogenated oil		Hunan cooking	Cantonese cooking	Cantonese style	Sichuan style	Dongbei style	Hunan style	Western-style fast food cooking	Chinese cooking		
Units	µg/kg of meat cooked	µg/kg of vegetables cooked		µg/kg of vegetables cooked		µg/kg of potatoes cooked		ng/mg of particles emitted		(ng/mg of POM)				ng/mg of POM	
<i>n-alkanes</i>	gas+particle	gas	particle	gas	particle	gas	particle	particle	particle	particle	particle	particle	particle	particle	particle
n-Tetradecane	5,000	930		530	25	340	5	8.2	3.8	19±17	23±9	24±6	12±3	46±8	19
n-Pentadecane	6,240	690		660	35	430	37	18	13	70±11	75±11	93±13	46±9	146±48	71
n-Hexadecane	3,240	720	150	450	33	230	18	27.8	32.4	128±33	43±10	44±25	21±15	161±26	59
n-Heptadecane	400	170	34	650	95	310	33	58	45.7	139±44	50±12	45±13	29±6	234±61	66
n-Octadecane	1,450	77	28	180	16	96	11	55	34.1	71±24	55±19	57±33	23±6	137±77	51
n-Nonadecane	720	82	14	97	22	49	9	53.7	30.9	72±14	75±35	69±17	39±8	167±94	64
n-Eicosane	1,190	38	26	100	14	37	6	58.1	33.7	97±25	65±19	36±30	39±17	nd	59
n-Heneicosane	880	26	38	46	15	26	3	49.9	46.8	167±80	104±61	91±35	75±22	252±53	110
n-Docosane	610	19	10	36	10	20	1	42.8	49.2	192±64	143±76	34±41	65±59	73±118	109
n-Tricosane	560	25	18	36	5	19	1	54	54.5	214±71	161±71	80±13	91±26	394±67	137
n-Tetracosane	690	18	18	15	18	12		22.1	31.5	135±33	102±54	54±20	66±18	251±140	89
n-Pentacosane	780			7	13	9		22	29.1	292±102	303±105	102±89	205±46	549±124	225
n-Hexacosane	260		18		17			17.2	24.3	81±22	59±43	12±20	25±21	289±109	44
n-Heptacosane	650		340		470			13.2	32.1	181±60	116±41	59±19	74±13	366±135	107
n-Octacosane	1,140							11	17.1	62±69	45±69	nd	nd	242±114	27
n-Nonacosane	770							23.9	324.2	552±223	212±79	63±43	207±68	278±77	258
n-Triacontane								14.1	19.2	16±25	31±23	nd	nd	131±66	12
n-Hentriacontane								47.9	99	560±197	367±149	96±51	346±45	121±44	342
n-Dotriacontane								ND	10.1					28±67	Nd
n-Tritriacontane								ND	19.7	26±41	90±55	20±32	nd	nd	34
Total class								596.9	950.3					3863	1883

Table 12. Fatty acids emitted from cooking food - indoor concentrations and emission factors (see columns for units)

Reference	Schauer et al., 1999a	Schauer et al., 2002						He et al., 2004b		Zhao et al., 2007b			
Compound class	Burger meat charbroiled	stir fry in soybean oil		stir fry in canola oil		deep frying of potatoes in hydrogenated oil		Hunan cooking	Cantonese cooking	Cantonese style	Sichuan style	Dongbei style	Hunan style
Units	µg/kg of meat cooked	µg/kg of vegetables cooked		µg/kg of vegetables cooked		µg/kg of potatoes cooked		ng/mg of particles emitted		ng/mg of POM			
Compound class	gas+particle	gas	particle	gas	particle	gas	particle	particle	particle	particle	particle	particle	particle
<i>n - alkanolic acids</i>													
Hexanoic acid								1,999	1,337	85 ± 74	159 ± 45	144 ± 112	103 ± 39
Heptanoic acid	32,200	5,940		530	162	370	2	2,295	1,688	13 ± 20	59 ± 11	67 ± 29	50 ± 18
Octanoic acid	38,700	5,930	160	4,330	170	640	27	3,202	2,444	32 ± 32	130 ± 32	137 ± 87	96 ± 27
Nonanoic acid	48,400	11,890	180	12,200	180	3,270	51	850	687	60 ± 46	208 ± 35	417 ± 243	154 ± 38
Decanoic acid	11,100	780	28	700		190	9	593	246	14 ± 16	60 ± 18	81 ± 27	44 ± 14
Undecanoic acid								390	90	3 ± 7	37 ± 11	73 ± 26	30 ± 7
Dodecanoic acid	6,460	320	59	210	20	100	4	2,462	463	116 ± 60	247 ± 86	208 ± 79	305 ± 113
Tridecanoic acid	21,700							422	92	25 ± 28	108 ± 47	113 ± 34	61 ± 15
Tetradecanoic acid		130	93	87	42	60	52	7,359	1,211	341 ± 114	1635 ± 809	801 ± 113	730 ± 188
Pentadecanoic acid	5,970		58		59		12	774	316	102 ± 27	437 ± 200	283 ± 43	224 ± 80
Hexadecanoic acid	17,400	238	2,980	690	2,280	800	1,760	57,892	26,621	10,608 ± 2,238	30,121 ± 7,982	23,344 ± 3,547	14,757 ± 2,947
Heptadecanoic acid	10,300				50		23	839	464	118 ± 41	285 ± 121	151 ± 28	162 ± 37
Octadecanoic acid	96,100	15	1,250	65	1,040	37	848	21,412	11,166	4,209 ± 996	6,084 ± 1,608	2,876 ± 533	3,975 ± 799
Nonadecanoic acid	600				7			47	50	5 ± 8	43 ± 10	29 ± 16	23 ± 7
Eicosanoic acid	860		38		65		48	1,959	2,462	257 ± 74	400 ± 119	191 ± 33	235 ± 54
Heneicosanoic acid								46	71				
Docosanoic acid	350		29		35		7	677	1,229	355 ± 94	490 ± 189	52 ± 52	268 ± 55
Tricosanoic acid								81	82				
Tetracosanoic acid								269	431	22 ± 35	86 ± 80	10 ± 24	71 ± 29
Pentacosanoic acid								23	20				
Hexacosanoic acid								22	29				
<i>n - alkenolic acids</i>													
9-hexadecenoic acid (palmitoleic acid)	18,400		174		36		18	3,638	2,101	101 ± 39	255 ± 96	108 ± 37	196 ± 60
9,12-octadecadienoic acid (Linoleic acid)	214,000		4,190		3,030		1,750	85,635	64,756	8,677 ± 2,233	13,547 ± 5,345	3,077 ± 904	10,132 ± 2,498
9-octadecenoic acid (oleic acid)	32,000		3,250		6,310		1,940	33,584	39,806	13,775 ± 2,663	29,375 ± 8,307	18,828 ± 3,256	18,044 ± 2,723
Eicosenoic acid								782	3,435				
13-Docosenoic acid								307	670				
9,12,15-octadecatrienoic acid (linolenic acid)			310		270		77						

Table 13. Dicarboxylic acids emitted from cooking food - indoor concentrations and emission factors (see columns for units)

Reference	Schauer et al., 1999a	Schauer et al., 2002			He et al., 2004b		Zhao et al., 2007b			
Compound class	Hamburger meat charbroiled	stir fry in soybean oil	stir fry in canola oil	deep frying of potatoes in hydrogenated oil	Hunan cooking	Cantonese cooking	Cantonese style	Sichuan style	Dongbei style	Hunan style
Units	µg/kg of meat cooked	µg/kg of vegetables cooked	µg/kg of vegetables cooked	µg/kg of potatoes cooked	ng/mg of particles emitted		ng/mg of POM			
butanedioic acid (Succinic acid)							49 ± 47	225 ± 64	262 ± 159	166 ± 43
pentanedioic acid (Glutaric acid)					667	687	4 ± 7	80 ± 28	91 ± 42	56 ± 16
hexanedioic acid (Adipic acid)	1,990		33		429	844	15 ± 17	129 ± 38	109 ± 31	70 ± 13
heptanedioic acid (Pimelic acid)					370	356	84 ± 42	246 ± 65	197 ± 73	154 ± 57
octanedioic acid (Suberic acid)	3,900	58	165	3	1,193	1,326	131 ± 75	413 ± 177	189 ± 72	201 ± 61
nonanedioic acid (Azelaic acid)					3,160	4,934	675 ± 284	1,890 ± 828	1,043 ± 671	975 ± 325
decanedioic acid (Sebacic acid)					403	350	97 ± 21	179 ± 62	156 ± 43	107 ± 29
undecanedioic acid							14 ± 18	82 ± 43	62 ± 24	50 ± 21

Table 14. Aldehydes emitted from cooking food - indoor concentrations and emission factors (see columns for units)

Reference	Schauer et al., 1999a	Schauer et al., 2002			Svendsen et al., 2002				He et al., 2004b				Zhao et al., 2007c		Zhao et al., 2007b				Sjaastad and Svendsen, 2008				Sjaastad and Svendsen, 2009
Aldehydes	Hamburger meat charbroiled	stir fry in soybean oil	stir fry in canola oil	deep frying potatoes in hydrogenated oil	Hotel kitchen	Burger chain	Restaurant with grill	Small local restaurants	Hunan cooking	Cantonese cooking	Hunan cooking	Cantonese cooking	Western-style fastfood	Chinese cooking (average)	Cantonese style	Schuan style	Dongbei style	Hunan style	Margarine	Rapeseed oil	Soybean oil	Olive oil	Norwegian à la carte restaurant
Units	µg/kg of meat cooked	µg/kg of vegetables	µg/kg of vegetables	µg/kg of potatoes	µg/m ³				ng/mg of particles emitted		ng/m ³		ng/mg of POM				µg/m ³						
formaldehyde	1,382,000	20,100	18,600	12,400	11 ± 8	7 ± 0	15 ± 11	14 ± 6															
acetaldehyde	1,092,000	50,100	42,200	20,900	29 ± 30	16 ± 5	34 ± 24	102 ± 33															
propanal	504,000	12,200	17,000	7,000																			
butanal/isobutanal	373,000	19,700	17,400	4,500																			
hexanal	203,000	4,100	6,400	6,700																			
heptanal	125,000	4,300	8,000	5,200																			
octanal	146,000	7,900	9,700	5,700																			
nonanal	148,000	12,400	14,800	13,500					973	827	1,369	555	25,518±7951	2380	1365 ± 393	2281 ± 443	4501 ± 2735	1373 ± 519					
decanal	33,600	5,200	1,090	2,900									651±244	225	49 ± 81	272 ± 80	405 ± 86	174 ± 34					
undecanal	17,200	3,000	200	1,200									430±180	138	33 ± 82	154 ± 92	255 ± 46	109 ± 25					
dodecanal	30,000	1,260	920										352±108	140	8 ± 21	147 ± 100	287 ± 83	120 ± 46					
tridecanal	18,000	550	180										514±127	102	nd	117 ± 72	193 ± 54	99 ± 37					
tetradecanal	18,000	410											536±154	199	62 ± 69	235 ± 59	359 ± 50	141 ± 39					
pentadecanal	14,400	440	411										1172±389	259	62 ± 69	305 ± 123	436 ± 104	232 ± 75					
Acrolein					11 ± 10	14 ± 9	12 ± 11	3 ± 1															
trans,trans-2,4-decadienal																			10.33 (2.52)	0.63 (1.32)	0.52 (0.8)	n.d.	9.79 (14.44)
2,4-Decadienal																			25.33 (4.51)	n.d.	n.d.	n.d.	3.12 (4.06)
trans-2-decenal																			25.33 (9.70)	3.60 (6.40)	0.50 (1.20)	0.50 (1.20)	8.23 (9.27)
s-2-decenal		16,100	26,400																n.d.	0.82 (1.08)	2.20 (5.29)	3.67 (2.94)	
2-Undecenal		18,400	29,400																20.67 (7.64)	3.81 (5.21)	2.02 (3.62)	3.33 (2.34)	8.23 (10.07)

Table 15. Ketones and lactones emitted from cooking food - indoor concentrations and emission factors (see columns for units)

Reference	Schauer et al., 2002				McDonald et al., 2003						Zhao et al., 2007b				Zhao et al., 2007c			
Compound class	stir fry in soybean oil		stir fry in canola oil		deep frying of potatoes in hydrogenated oil		Hamburger Auto-Char	Hamburger Under-Char	Steak Under-Char	Chicken Under-Char	Hamburger Griddle	Chicken Griddle	Cantonese style	Sichuan style	Dongbei style	Hunan style	Western-style fast food	Chinese cooking
Units	µg/kg of vegetables cooked				µg/kg of potatoes		mg/kg						ng/mg of POM					
Compound class	Gas	Particle	Gas	Particle	Gas	Particle	Particle	Particle	Particle	Particle	Particle	Particle	Particle	Particle	Particle	Particle	Particle	Particle
Ketones																		
2-nonanone	3300				78		2.69	11.95	7.50	11.35	0.55	0.59	116±90	326±90	504±44	213±43	240±99	290
2-decanone	2670		3130		590		1.27	5.74	4.06	4.46	0.26	0.58	116±91	361±108	589±45	235±46	203±165	325
2-undecanone	2310				145		1.03	4.25	3.52	2.73	0.28	0.26	58±50	259±72	386±39	197±44	110±135	225
2-Dodecanone													nd	102±42	151±19	69±12	10±24	81
2-tridecanone					84								nd	35±86	64±100	20±49	17±40	30
2-tetradecanone					180								nd	23±56	103±54	65±95	28±69	48
2-pentadecanone	3900	170	8050	120	1100	30							234±152	592±262	844±181	401±138	5419±1288	518
2-Hexadecanone													nd	16±26	39±51	8±19	994±177	16
2-heptadecanone	720	60	860	75	300								667±180	1003±489	1383±330	594±134	15,682±2321	912
Furanones (Lactones)																		
5-ethylidihydro-2(3H)-furanone	470		370		41													
5-propylidihydro-2(3H)-furanone	170		170		11								6±16	70±41	86±35	53±17	24±29	54
5-butylidihydro-2(3H)-furanone	430	17	670	30	240	10							21±45	188±96	172±47	177±67	211±59	140
5-pentylidihydro-2(3H)-furanone	280	30	470	38	75	7	2.69	11.95	7.5	11.35	0.55	0.59	20±42	221±138	288±135	134±44	208±76	166
5-hexylidihydro-2(3H)-furanone	74	45	130	40	49	9	1.27	5.74	4.06	4.46	0.26	0.58	8±21	90±44	119±39	56±15	109±45	69
5-heptylidihydro-2(3H)-furanone	33	5		5		3							5±11	65±28	102±34	39±9	94±18	53
5-octylidihydro-2(3H)-furanone		43		53		1							12±20	74±37	120±35	94±33	228±65	75
5-nonylidihydro-2(3H)-furanone		29		33		2							4±9	71±24	91±11	37±9	358±69	51
5-decylidihydro-2(3H)-furanone		3		4		1							43±40	117±46	157±35	58±30	774±604	94
5-undecylidihydro-2(3H)-furanone		41		30		2	1.03	4.25	3.52	2.73	0.28	0.26	50±56	179±74	147±42	84±14	1212±337	115
5-dodecylidihydro-2(3H)-furanone		12		10		1							809±125	1401±466	1260±182	739±135	6646±1235	1052
5-Tridecylidihydro-2(3H)-furanone													19±23	58±22	7±11	15±13	885±180	25
5-Tetradecylidihydro-2(3H)-furanone													329±78	311±112	178±35	184±37	2573±661	250

Table 16. Concentrations of organic ions emitted from cooking food

Reference	See and Balasubramanian, 2008				
Pollutant	Steaming	Boiling	Stir-frying	Pan-frying	Deep-frying
Units	$\mu\text{g}/\text{m}^3$	$\mu\text{g}/\text{m}^3$	$\mu\text{g}/\text{m}^3$	$\mu\text{g}/\text{m}^3$	$\mu\text{g}/\text{m}^3$
Organic Ions					
Acetate	0.09 ± 0.02	0.19 ± 0.02	0.67 ± 0.05	0.77 ± 0.07	1.0 ± 0.1
Formate	0.07 ± 0.01	0.14 ± 0.02	0.32 ± 0.03	0.33 ± 0.04	0.39 ± 0.05
Methanesulfonate	BDL	BDL	BDL	BDL	BDL
Pyruvate	BDL	BDL	BDL	BDL	BDL
Succinate	2.2 ± 0.4	3.3 ± 0.6	5.0 ± 0.8	7.8 ± 0.9	14.3 ± 1.6
Glutarate	0.01 ± 0.00	0.01 ± 0.00	0.02 ± 0.01	0.02 ± 0.01	0.04 ± 0.01
Malonate	BDL	BDL	BDL	BDL	BDL
Oxalate	0.39 ± 0.05	0.47 ± 0.06	0.78 ± 0.12	0.79 ± 0.15	0.93 ± 0.18

Table 17. Molecular markers, amides and other organic compounds emitted from cooking food - indoor concentrations and emission factors (see columns for units)

Reference	He et al., 2004b		Zhao et al., 2007b				Zhao et al., 2007c		McDonald et al., 2003					
Compound class	Hunan cooking	Cantonese cooking	Cantonese style	Sichuan style	Dongbei style	Hunan style	Western-style fast food cooking	Chinese cooking	Hamburger Auto-Char	Hamburger Under-Char	Steak Under-Char	Chicken Under-Char	Hamburger Griddle	Chicken Griddle
Units	ng/mg of particles emitted		ng/mg of POM				ng/mg of POM		mg/kg					
Sterols														
Cholesterol	525	369	261 ± 61	353 ± 125	114 ± 19	315 ± 84			1.35	7.44	1.41	7.87	0.004	0.29
Stigmasterol	336.9	145.2	619 ± 104	621 ± 245	84 ± 17	516 ± 133								
β -sitosterol	1352	2604	1168 ± 181	1313 ± 527	293 ± 31	1080 ± 286								
Monosaccharide Anhydrides														
Galactosan	2.2	6.1	nd	20 ± 7	30 ± 11	4 ± 4								
Mannosan	2.8	7.4	nd	7 ± 3	14 ± 9	3 ± 2								
Levogluosan	50.5	196.8	282 ± 147	218 ± 56	554 ± 296	124 ± 33								
Other compounds														
2-Pentylfuran	49.1	52.6	237 ± 127	237 ± 119	170 ± 228	82 ± 40								
Benzoic acid	ND	10	39 ± 11	30 ± 6	20 ± 6	17 ± 4								
Amides														
Tetradecanamide			nd	3 ± 8	3 ± 6	nd	841 ± 173	1						
Hexadecanamide			376 ± 99	249 ± 57	494 ± 137	172 ± 47	2797 ± 676	323						
9-Octadecenamide			185 ± 73	112 ± 23	165 ± 54	92 ± 23	344 ± 231	139						
Octadecanamide			129 ± 33	48 ± 25	53 ± 28	43 ± 13	710 ± 186	68						

Table 18. EC/OC, metals and inorganic ions emitted from cooking food -indoor concentrations and emission factors (see columns for units)

Reference	McDonald et al., 2003						See and Balasubramanian, 2008					See and Balasubramanian, 2006	He et al., 2004b	
Pollutant	Hamburger Auto-char	Hamburger Under-Char	Steak Under-Char	Chicken Under-Char	Hamburger Griddle	Chicken Griddle	Steaming	Boiling	Stir-frying	Pan-frying	Deep-frying	Chinese cooking in Commercial food stall	Hunan cooking	Cantonese cooking
Units	mg/Kg						µg/m ³					µg/m ³	wt%	
EC/OC														
EC	119	343.55	201.9	19.14	n.d.	n.d.	6.16 ± 0.71	8.11 ± 1.08	14.5 ± 1.4	14.7 ± 2.0	15.8 ± 2.2		1	1.6
OC	4464.21	15541.42	7061.25	6881.12	n.d.	n.d.	29.3 ± 3.6	36.1 ± 4.0	62.6 ± 5.6	71.6 ± 7.0	121.5 ± 16.3		81.6	52.6
Metals														
Al	0.95	4.88	0	0	n.d.	n.d.	61.7 ± 8.7	64.5 ± 9.4	72.3 ± 8.5	83.1 ± 8.1	137.3 ± 12.6	523.6±78.0		
As							10.5 ± 1.7	11.9 ± 1.9	17.9 ± 2.0	21.6 ± 2.6	28.9 ± 3.2	17.1±2.4		
Cd							2.17 ± 0.23	3.16 ± 0.52	6.67 ± 0.86	6.58 ± 0.68	13.1 ± 1.2	4.2±0.5		
Co							0.56 ± 0.07	0.70 ± 0.06	1.07 ± 0.11	1.61 ± 0.11	1.63 ± 0.24	9.7±1.0		
Cr							29.7 ± 2.9	30.4 ± 2.7	57.0 ± 4.0	68.4 ± 5.3	95.2 ± 9.2	137.1±14.6		
Cu	0.13	0	0	0	n.d.	n.d.	367.2 ± 20.9	416.5 ± 41.7	670.1 ± 54.9	1093 ± 126	1107 ± 119	3534.5±426.2		
Fe	1.76	6.06	0	0	n.d.	n.d.	441.5 ± 59.5	527.4 ± 68.0	1693 ± 196	3157 ± 335	4527 ± 463	4754.8±480.9		
Mn							12.5 ± 1.3	19.5 ± 1.9	28.9 ± 3.0	54.5 ± 4.6	62.8 ± 8.5	128.6±14.9		
Ni							20.9 ± 2.1	24.0 ± 4.2	30.7 ± 3.1	45.2 ± 3.2	71.2 ± 10.2	119.8±14.1		
Pb							10.6 ± 1.4	12.2 ± 1.3	52.5 ± 4.6	55.7 ± 6.2	109.6 ± 12.5	480.0±50.8		
V							16.7 ± 2.3	20.5 ± 2.3	22.9 ± 2.4	30.8 ± 2.9	74.1 ± 8.6	272.2±29.0		
Ti							69.0 ± 8.6	102.5 ± 14.4	190.0 ± 27.1	223.9 ± 22.4	368.8 ± 51.6			
Zn	4.15	0	0	0	n.d.	n.d.	529.9 ± 48.8	558.0 ± 38.9	796.0 ± 53.8	815.0 ± 73.8	937.4 ± 110.6	5486.9±755.7		
Sb												163.0±16.6		
Inorganic Ions														
Li ⁺							BDL	BDL	BDL	BDL	BDL			
Na ⁺							0.48 ± 0.08	0.54 ± 0.09	0.44 ± 0.05	0.45 ± 0.08	0.56 ± 0.11		0.29	0.15
NH ₄ ⁺	0.52	2.53	0	0	n.d.	n.d.	1.0 ± 0.1	1.2 ± 0.1	1.1 ± 0.1	1.3 ± 0.1	1.3 ± 0.3		0.13	0.34
K ⁺	12.81	60.14	4.86	0	n.d.	n.d.	1.2 ± 0.1	1.3 ± 0.2	0.87 ± 0.11	1.0 ± 0.2	1.1 ± 0.3		0.06	0.05
Mg ²⁺							BDL	BDL	BDL	BDL	BDL		0.02	0.02
Ca ²⁺							0.63 ± 0.08	0.73 ± 0.10	0.59 ± 0.08	0.95 ± 0.09	0.99 ± 0.10		0.1	0.1
F ⁻							1.9 ± 0.4	3.2 ± 0.5	0.52 ± 0.07	0.55 ± 0.08	1.1 ± 0.2			
Cl ⁻	3.25	14.23	0	0	n.d.	n.d.	1.3 ± 0.2	2.6 ± 0.3	0.39 ± 0.04	0.56 ± 0.08	0.98 ± 0.11		0.3	0.24
NO ₂ ⁻							BDL	BDL	BDL	BDL	BDL			
NO ₃ ⁻	0.54	7.15	0	0	n.d.	n.d.	5.1 ± 0.6	6.0 ± 0.6	4.4 ± 0.4	4.5 ± 0.4	5.0 ± 0.4		0.32	0.34
SO ₄ ²⁻	4.68	17.01	0	0	n.d.	n.d.	3.9 ± 0.5	4.1 ± 0.7	3.1 ± 0.3	3.3 ± 0.4	3.4 ± 0.5		0.23	0.77
PO ₄ ³⁻							BDL	BDL	BDL	BDL	BDL			

Table 19. Comparison of diagnostic ratios of PAHs

REFERENCES	SOURCE OF PM	Pollutant	Phe/ (Phe+Ant)	Flu/ (Flu+Pyr)	BaA/ (BaA+Chry)	Ind/ (Ind+Bpe)	BeP/ (BaP+BeP)	Ant/ (Ant+Phe)	IcP/ (IcP+BgP)
COOKING STYLES									
See et al., 2006	CHINESE	PM _{2.5}	0.21	0.32	0.4	0.43			
	MALAY	PM _{2.5}	0.28	0.38	0.32	0.38			
	INDIAN	PM _{2.5}	0.21	0.43	0.5	0.39			
Li et al., 2003	CHINESE	TSP and gas	0.86	0.5	0.62	0.63			
	WESTERN	TSP and gas	0.86	0.46	0.38	0.63			
	FAST FOOD	TSP and gas	0.96	0.6	0.32	0.53			
	JAPANESE	TSP and gas	0.97	0.66	0.13	0.83			
He et al., 2004	CHINESE, HUNAN	PM _{2.5}	0.96	0.44	0.51	–			
	CHINESE, CANTONESE	PM _{2.5}	1	0.36	0.47	0.19			
Zhu and Wang, 2003	CHINESE	TSP and gas	0.51	0.18	0.74	–			
	CHINESE	TSP and gas	0.41	0.19	0.18	–			
	CHINESE	TSP and gas	0.37	0.23	0.22	–			
	CHINESE	TSP and gas	0.51	0.23	0.38	–			
COOKING METHODS									
See and Balasubramanian, 2008	STEAMING	PM _{2.5}	0.96	0.51	0.31	0.54			
	BOILING	PM _{2.5}	0.96	0.52	0.34	0.52			
	STIR-FRYING	PM _{2.5}	0.94	0.56	0.28	0.45			
	PAN-FRYING	PM _{2.5}	0.94	0.55	0.29	0.46			
	DEEP-FRYING	PM _{2.5}	0.95	0.53	0.28	0.44			
OTHER SOURCES									
CITY OF SHANGAI Gu et al., 2010	TRAFFIC			0.52	0.27		0.63	0.13	0.45
Miguel and Pereira, 1989	TRAFFIC			0.4-.5					
	PETROLEUM			<0.2					
	GRASS			>0.5					

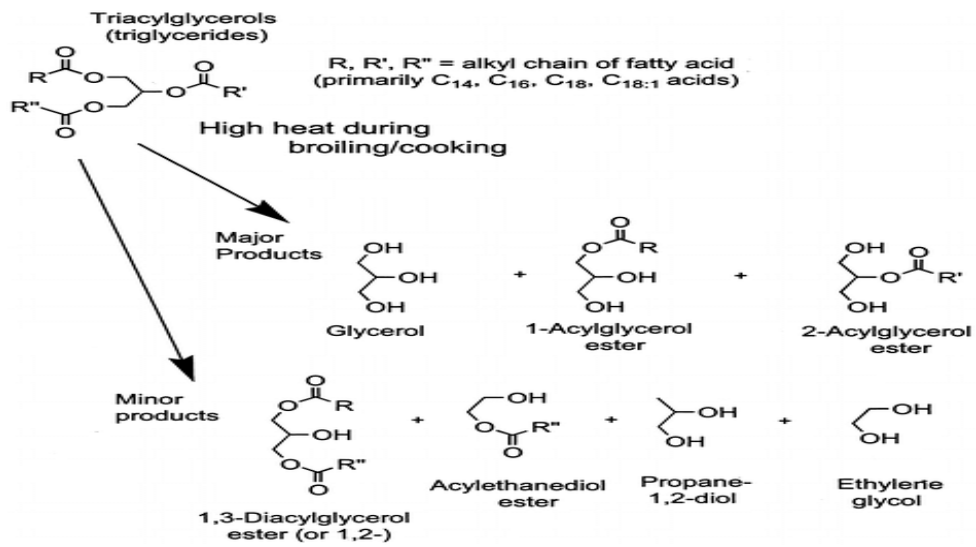


Figure 1: Break down products of triglycerides (Nolte et al., 1999)

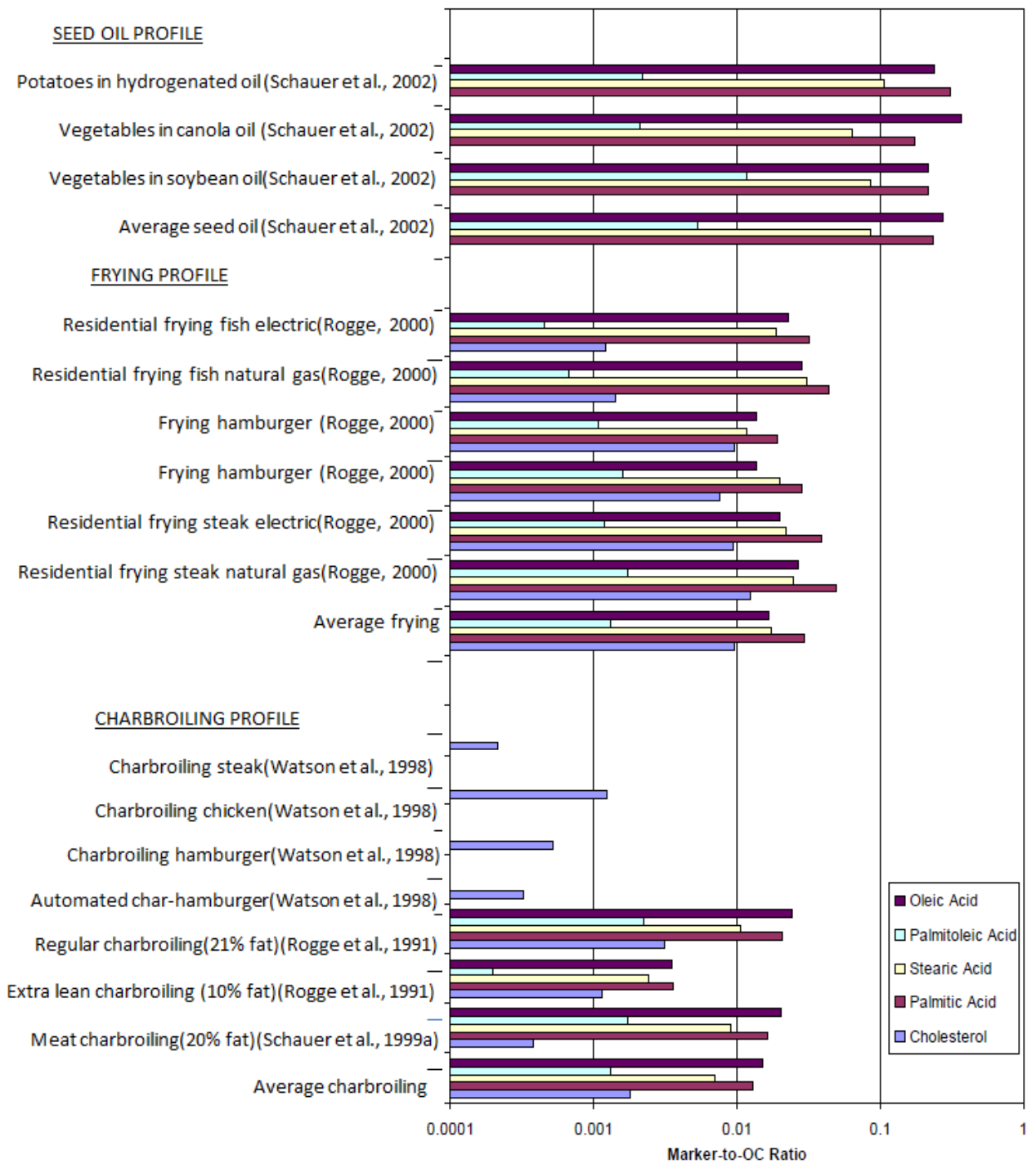


Figure 2: Marker to OC ratio for meat cooking profiles (Robinson et al., 2006)