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A Further Tool To Monitor Coffee Roasting Process: Aroma Composition And Chemical Indices

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1 Non-separative headspace solid phase microextraction-mass spectrometry profile as a marker
2 to monitor coffee roasting degree

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8

9 **Abstract**

10 This study describes a non-separative Headspace-Solid-Phase-Microextraction-Mass-
11 Spectrometry (HS-SPME-MS) approach, in view of its application to on-line
12 monitoring of a roasting process. The system can quickly provide representative and
13 diagnostic fingerprints of the volatile fraction of samples and, in combination with
14 appropriate chemometric pattern-recognition and regression techniques, can
15 successfully be applied to characterize, discriminate and/or correlate patterns with the
16 roasting process. Eighty coffee samples of different varieties, geographical origins
17 and blends were analyzed. The experimental HS-SPME-MS results show that the TIC
18 fingerprint can be used to discriminate the degree of roasting; diagnostic ion
19 abundance(s) or ratios were closely correlated with the roasting process; both could
20 successfully be used as markers or Analytical Decision Makers, to monitor roasting
21 processes on-line, and to define quality and safety of roasted coffee.

22
23 **KEYWORDS:** HS-SPME-MS; MS-EN; coffee; aroma quality and safety;
24 technological process; roasting indices; on-line monitoring
25

26 INTRODUCTION

27 The roasting process is a significant factor in determining coffee flavor. In particular,
28 coffee aroma depends on the specific quali-quantitative distribution of various
29 components. These are mainly volatile medium-to-high polarity compounds, deriving
30 from drying and from the heat-browning related to the roasting conditions, above all
31 temperature and time. The aroma also depends on the species, variety and blend, as
32 well as on geographical origin¹⁻⁴. Conversely, the roasting process can also produce
33 some compounds that must be monitored because of their toxic properties; this is the
34 case of furan and its derivatives, which have shown carcinogenic and cytotoxic
35 activities⁵⁻⁸. The reduction of furan in coffee is therefore highly recommended, and
36 may be achieved by optimizing the roasting process in all its steps (i.e. roasting,
37 cooling, degassing and grinding) while, obviously, keeping the organoleptic
38 properties unaltered.

39 The effect of the roasting process on coffee beans is generally described in terms of
40 the degree of roasting, and is usually evaluated through several chemical and physical
41 parameters, including external color of the beans, loss of weight during roasting, and
42 variations in chemical composition, as well as through the development of sensory
43 characteristics⁹⁻¹⁰. These parameters concur to define the degree of roasting, although
44 to date a concise, clear and universally-accepted evaluation protocol does not exist.
45 The most widely-used parameter to determine degree of roasting in day-to-day
46 practice is color, measured through the light reflectance of ground beans or, still
47 today, by visual inspection. This latter method is still valid partly because the
48 industry uses a constant quality and variety of green coffee, combined with constant
49 time-temperature conditions and roasting plant. Dry-matter loss is also considered to
50 provide a reliable and complementary evaluation of the degree of roasting, including
51 for in-plant determination⁹. Correct conditions for industrial roasting processes are, in
52 general, obtained by appropriately scaling-up the results of experiments from pilot or
53 laboratory plants monitored by physical methods.

54 This study describes a non-separative headspace solid phase microextraction-mass
55 spectrometric method (HS-SPME-MS), in view of its possible application to the on-
56 line monitoring of roasting processes. HS-SPME is a high-concentration-capacity
57 technique offering good recoveries and easy to automate, that can be combined
58 directly on-line with mass spectrometry¹¹. Non-separative MS methods, better known
59 as mass spectrometry-based electronic nose (MS-EN)¹²⁻¹⁶, were introduced by
60 Marsili¹⁷ to study off-flavors in milk; they have since been applied successfully to
61 characterizing several matrices, in particular in the food field¹⁸⁻²⁰. They provide a
62 representative, diagnostic and generalized mass spectrometric fingerprint of the
63 volatile fraction of a sample, analyzed directly without prior chromatographic
64 separation, in which each m/z ratio acts as a “sensor” whose intensity derives from
65 the contribution of each compound producing that fragment. These methods, in
66 combination with appropriate chemometric elaboration can be used to quickly
67 characterize and discriminate samples within a set, and to correlate them with a
68 technological process (e.g. coffee roasting). MS-EN can also be used to monitor

69 target compounds in a group of samples, provided that specific and diagnostic ions
70 are obtained with a compatible ion generation mode (EI, CI, APCI, PTR etc.).
71 Mass spectral fingerprints, or diagnostic ion abundance(s), were here used both as
72 marker and as analytical decision maker (ADM)²¹ to monitor coffee roasting degree,
73 in view of the possibility of combining a mass spectrometer directly on-line to a
74 roasting machine. In particular, the results are reported of a study aimed at correlating
75 HS-SPME-MS profile and coffee color, as a further tool to characterize roasting
76 processes.

77

78 **MATERIALS AND METHODS**

79 ***Samples:***

80 Eighty roasted-coffee samples from different geographical origins, of different
81 varieties and blends (100% Arabica, 100% Robusta, 50/50% Arabica-Robusta blend,
82 and several commercial blends) at different degrees of roasting were supplied by
83 Lavazza (Turin – Italy) over a period of 12 months. Colors of samples were
84 measured, by ground-bean light reflectance, with a single-beam Neuhaus Neotec
85 Color Test II instrument, at wavelength 900 nm, on 25-30g of ground coffee. Table 1
86 gives the list of samples with their color values. A set of 100 coffee pods from the
87 same lot (a 50/50 Arabica and Robusta blend) for espresso machines was stored at -
88 18°C and used as reference to standardize the HS-SPME system performance over
89 time.

90

91 ***Headspace Solid Phase Microextraction (HS-SPME) sampling and GC analysis***
92 ***conditions***

93 The SPME device and fibers were from Supelco (Bellefonte, PA, USA).
94 Divinylbenzene/Carboxen/ Polydimethylsiloxane (DVB/CAR/PDMS) d_f 50/30 μm , 2
95 cm long fibers were used, conditioning them before use as recommended by the
96 manufacturer²².

97 Volatiles were sampled by automated headspace Solid Phase Microextraction (auto-
98 HS-SPME) using a MPS-2 multipurpose sampler (Gerstel, Mülheim a/d Ruhr,
99 Germany) on-line integrated with an Agilent 7890 GC unit coupled to an Agilent
100 5975 MS detector (Agilent, Little Falls, DE, USA). 500 mg of ground roasted coffee
101 in a 20mL screw-cap vial were sampled by HS-SPME at 50°C for 10 minutes¹⁰. All
102 samples were analyzed in triplicate. Three fibers were selected from different lots,
103 after preliminary testing to establish their sampling performance, so as to select
104 equivalent fibers for use throughout the entire analysis period (see below: HS-SPME
105 fiber performance evaluation). Fiber performance was monitored throughout the
106 study on an additional set of coffee pod samples, and on 5 μL of a 2 mg/mL standard
107 solution of α - and β -thujone in dibutyl phthalate.

108 *Non-separative analysis – GC unit conditions:* oven and injector temperature: 250 °C
109 injection mode: split, split ratio: 1/10; carrier gas: helium, flow rate: 0.4 mL/min;
110 fiber desorption time and reconditioning: 3min; transfer column: deactivated fused
111 silica tubing (d_c 0.10mm, length 6.70m) (Mega, Legnano (Milan), Italy).

112 *MSD conditions:* ionization: EI mode at 70 eV; transfer line: 280°C. Standard tuning
113 was used and the scan range was set at m/z 35-350 with a scanning rate of 1.000
114 amu/s.

115 *Separative GC-MS analysis - Chromatographic conditions:* injector temperature:
116 230°C, injection mode: split, ratio: 1/10; carrier gas: helium, flow rate: 1 mL/min;
117 fiber desorption time and reconditioning: 5 min; column: MEGAWAX 20M (d_f 0.20
118 μm , d_c 0.20 mm, length 50 m) (Mega, Legnano (Milan), Italy). Temperature program:
119 from 40°C (1 min) to 200°C at 3°C/min, then to 250 °C (5min) at 10°C/min.

120 *MSD conditions:* ionization mode: EI (70 eV), scan range: 35 to 350 amu; ion source
121 temperature: 230°C; quadrupole temperature: 150°C; transfer line temperature:
122 250°C.

123 Data were acquired and processed with an Agilent MSD Chem Station ver
124 E.02.01.1177. (Agilent, Little Falls, DE, USA)

125 Analytes were identified by their linear retention indices and EI-MS spectra, by
126 comparison with authentic standards, or were tentatively identified through their EI-
127 MS fragmentation patterns and retention indices.

128
129 ***Chemometric analyses***

130 Principal Component Analysis (PCA) and Orthogonal Partial Least Square analysis
131 (OPLS) were performed with Pirouette software ver. 4.0 (Infometrix, Inc. Bothell,
132 WA). The software package was used to automatically create ASCII files from
133 Agilent GC ChemStation data, using a post-run macro. The data matrix consisted of
134 as many rows as the number of samples (total objects: 240) and 316 columns (m/z

135 variables). Samples were randomly divided into a training set (55 samples) and a test
136 set (25 samples). PCA was used in the first step for pattern recognition analysis, to
137 visualize information and sample clusters, in particular as a function of the
138 technological process. OPLS analysis was then carried out to correlate color (as a
139 marker of the degree of roasting) to the chemical fingerprint. Data were pre-treated
140 by baseline correction, through noise subtraction, and by internal normalization of the
141 signal from each sample; they were subsequently pre-processed by mean-centering.
142

143 **RESULTS AND DISCUSSION**

144 Compared to separative GC-MS profiles, MS-EN patterns provide a fast response;
145 however, the significance of the TIC is low, and less information can apparently be
146 obtained from the MS profile. This is because the intensity of each fragment (m/z) is
147 “built up” from the contributions of each component of the sample involved in
148 generating that ion during its ionization process. Figure 1 gives the HS-SPME-GC-
149 MS (a) and the HS-SPME-MS TIC (b) profiles, together with the mass spectral
150 fingerprint (c) of an Arabica coffee sample. Table 2 lists the components identified in
151 the HS-SPME-GC-MS profile of the same Arabica sample, together with their target
152 (TI) and qualifier ions (Qi).

153 When applying HS-SPME-MS, it is mandatory to use chemometric techniques to
154 “extract” data from the MS profile that can provide significant information. However,
155 when a mathematical model is generated and used to classify or correlate
156 composition or characteristics of a sample, and to reveal the resources of hidden
157 information, one of the main limits is profile precision over a long period. This
158 variability can reduce the effectiveness of a successful model, generated with a
159 training set, when applied to routine analyses. The most critical points in the case of
160 HS-SPME-MS are SPME fiber performance and MS signal instability; the latter may
161 be due to ion source contamination, aging of electron multiplier, and/or filament
162 electron emission. These limits can be overcome by i) evaluating the response of a
163 suitable number of SPME fibers and monitoring it continually, ii) checking electron
164 multiplier response over time, and iii) standardizing the MS profile through internal
165 normalization.
166

167 **HS-SPME fiber performance evaluation**

168 SPME fiber effectiveness was evaluated initially in terms of total fingerprint areas:
169 this was aimed at minimizing sampling errors/discriminations, in view of the
170 extended time interval (12 months) covered by this study, and of the large number of
171 samples and replicates expected to be run. As discussed elsewhere²¹, the sampling
172 performance of three DVB/CAR/PDMS fibers from different lots was tested on
173 coffee pod samples and on an α - and β -thujone standard solution, to classify them on
174 the basis of the recovery provided, and to monitor any change over time. Normalized
175 spectral fingerprint areas of the coffee volatile fraction of five replicates from the
176 same pod for each fiber (F1, F2, F3) were submitted to Analysis of Variance
177 (ANOVA). The One-Way ANOVA results on the replicates for each fiber are in table
178 S1 (supporting information), and showed that the null hypothesis (“there is no

179 difference among the fibers in terms of absolute peak areas of the selected target
180 analytes“) was false. Tukey’s test classified F1 and F3 as belonging to the same
181 group, and these were therefore adopted for all experiments. Similar results were
182 obtained with α - and β -thujone standard solution. When additional fibers were
183 necessary, they were submitted to the entire test routine, analyzing reference coffee-
184 pod samples. Normalized spectral fingerprint area values had to fall within 5%
185 variability (expressed as RSD%) as established during performance testing.

186 187 **Precision and internal normalization**

188 Precision, expressed as repeatability and intermediate precision of HS-SPME-MS on
189 reference coffee-pod samples, was evaluated over the entire period. Repeatability was
190 calculated over five analyses of coffee in the same pod, the MS profile fragments of
191 each sample were normalized *versus* the most intense ions (m/z 43, basic peak) taken
192 as 1; each m/z intensity value is expressed as a percentage of the intensity of the basic
193 m/z fragment^{14,24}. Repeatability is expressed as Relative Standard Deviation %
194 (RSD%) on the total areas of the normalized fingerprints, and on some diagnostic m/z
195 ions characteristic of certain components related to roasting, aroma or toxic
196 chemicals. Intermediate precision ~~was calculated as before~~ was calculated as
197 ~~described above?// was calculated as described elsewhere?~~ over five analyses of the
198 coffee pod, carried out monthly over a period of one year. Results are in table S2
199 (supporting information) and show very good intermediate precision, with RSD%
200 values for the total fingerprinting area of 3.16%, and for single ions ranging from
201 4.65% for m/z 45 to 20.08% for m/z 150.

202 203 **Unsupervised exploratory analysis**

204 Although color is the most widely-adopted parameter used to monitor the roasting
205 process, on-line monitoring of the development of the volatile fraction by means of
206 an analytical approach (HS-SPME-MS) could be very useful to improve coffee
207 standardization and to optimize its aroma. Fifty five coffee samples of different
208 varieties and blends, and roasted to different degrees, i.e. dark (color range: 35÷42),
209 medium (46÷53) and light (57÷62), were analyzed. Their spectral fingerprints were
210 processed using an unsupervised approach, Principal Component Analysis (PCA),
211 with the aim of finding inter-sample and inter-variable relationships with color (e.g.
212 degree of roasting). This was done by visualizing ~~spots? variables? scores?~~
213 ~~components?~~ sample distribution on the score plots resulting from exploratory
214 analysis³¹⁻³³. PCA clearly showed that coffee samples are discriminated by their
215 roasting degree (color) on the first principal component (1st PC), while different
216 varieties are separated on the 2nd and 3rd PCs (Figure 2a). The volatile fraction
217 included the well-known Maillard reaction products, derived from non-volatile
218 precursor degradation (Amadori compounds and deoxyosones) namely furanes,
219 furanones and pyranones. Compounds resulting from the Strecker reaction of α -
220 dicarbonyls and aminoacids were also present: 2,3-butandione, 2- and 3-
221 methylbutanal and alkyl-pyrazines, as were sugar degradation products (furfural, 5-
222 methylfurfural and furfuryl alcohol). The loading plot (figure 2b) indicates those m/z

223 fragments that vary linearly with roasting degree; this indirectly also shows which
224 components change during roasting: m/z 52 and 79 (mainly representative of pyridine
225 and furfuryl acetate), 81 (1-methyl-pyrrole, 1-furfuryl-pyrrole and furfuryl acetate),
226 98 (furfuryl alcohol), 108 (dimethyl pyrazine groups: 2,3-, 2,5-, 2,6-isomers), 109
227 (guaiacol), 110 (5-methyl furfural), and 96 (furfural) all have high relevance ~~// are all~~
228 ~~highly significant?~~ on the 1st PC (explained variance 59.90%), and can be taken as
229 markers of roasting, as they increase markedly with the roasting degree. Some of
230 them are also key-aroma components, and are marked with an asterisk in Table
231 S2^{25,26}. M/z 43 (2,3-hexandione, acetoxyacetone), 45 (acetoine and 3-hydroxy-2-
232 butanone), 57 (2-oxopropyl propanoate), 60 (acetic acid), 79 (pyridine), and 95-96
233 (acetyl furan, furfural) on the 2nd PC (exp. var. 23.31%) are markers of variety and,
234 roasting degree ~~being equal????~~, are more abundant in Arabica than in the blend or
235 Robusta samples; they decrease with roasting ~~time/temperature~~. Conversely, m/z 108
236 (mainly deriving from dimethyl pyrazines) and 109 (guaiacol) are more intense in
237 Robusta samples. These considerations confirm the findings of a study by Ruosi et al.
238 using separative techniques.²⁷

239 Some m/z variables vary linearly with roasting degree, for example furan and its
240 homologues. Their formation is related linearly to the degree of roasting, as shown in
241 figure 3; determination coefficients were 0.7754 and 0.8917 for furan and 2-
242 methylfuran, respectively. Despite the presence of interference by the same m/z
243 fragment from other origins, this close correlation with degree of roasting means that
244 furan and 2-methyl furan formation can be monitored during roasting via their
245 characterizing ions (target ions), i.e. 68 and 82. If the intensities of these ions are
246 outside fixed limits of acceptance, the relative compounds can be quantified by
247 conventional methods^{28,29}. In addition to conventional separative methods, in 2011
248 Bicchi et al. proposed a quick method to quantify these compounds by HS-SPME-
249 MS; results were comparable to those of conventional methods³⁰.

250

251 **“Supervised” multivariate regression**

252 PCA results show that the MS profile of the coffee volatile fraction is closely
253 correlated to the degree of roasting, thanks to some variables (m/z values, and the
254 originating ~~// the relative?~~ components) that are characteristic of this technological
255 process. Orthogonal Partial Least Square analysis (OPLS) was thus used as a measure
256 of correlation, i.e. to evaluate the closeness of association between the fingerprint of
257 the volatile fraction of coffee and its color, as an indicator of technological treatment,
258 rather than for its ability to predict coffee color^{33,34}. The OPLS model requires a
259 training set to demonstrate any correlation between coffee color and aroma. Training
260 (55) and test (25) sets of samples, selected randomly and consisting of several
261 commercially-available Arabica (100%) and Robusta (100%) coffees of different
262 origins, plus blends of the two at different percentages, were established.

263 The OPLS regression model was first calculated on 5 PCs, and internally cross-
264 validated on the training set data relating to the volatile fraction spectral fingerprint
265 and coffee color; the model was then applied to the test set samples. The results
266 showed a highly negative linear correlation between measured (i.e. all m/z fragments

267 of the volatile fraction of coffee samples) and predicted (color values) variables. The
268 correlation between spectral fingerprint and color is negative due to how color values
269 are expressed, since high color values indicate lightly-roasted samples. ~~//In terms of~~
270 ~~predictive value, there was a close correlation (rpred 0.9472) with a satisfactory~~
271 ~~Standard Error of Prediction (SEP 2.53); values were within the range of colors that~~
272 ~~discriminate the different degree roasting (i.e. light, medium and dark).~~ TEMO DI
273 NON AVER FATTO GIUSTO, CI RIPROVO// _When used in prediction mode, the
274 correlation value was high (rpred 0.9472) with a satisfactory Standard Error of
275 Prediction (SEP 2.53); values were within the range of colors that discriminate the
276 different degree roasting (i.e. light, medium and dark).

277 Figure 4 shows the association intensity relationship between measured and predicted
278 color values, using the model built with the training set, and gives the parameters of
279 the validated model and of the prediction. As shown by PCA, the m/z ions with good
280 correlation values ($> \pm 0.7$) are related to compounds whose abundance is affected
281 markedly by the roasting process, and that are linked to the aroma developing during
282 this process (Fig 2b). The volatile fraction spectral fingerprint/color correlation is
283 important to monitor the roasting process because, besides establishing a relationship
284 between chemical and physical data, it ~~(sarebbe the correlation: OK?)~~ can give
285 specific chemical information about aroma changes with technological treatments.

286

287 **Identification of indices of the degree of roasting and possible relation with the** 288 **utility of roasted coffee color**

289 This part of the study investigated the possibility of identifying reliable chemical
290 indices to be used for roasting process control, as a function of some informative m/z
291 ion ratios. If correctly monitored on the laboratory scale, through objective and
292 measurable indices, the roasting process could be directed so as to obtain the required
293 aroma profile, in particular when new blends are being developed. Starting from the
294 above PCA results, the two-by-two normalized area ratios of ions closely correlated
295 to the roasting process were calculated, using a specific visual basic Excel macro.
296 The resulting indices were multiplied by 1000, to facilitate their handling, and were
297 arbitrarily considered significant only when there was a difference of at least 100
298 units between light, medium and dark roasting degree, in each variety or blend. Three
299 indices (i.e. m/z ratios) were found to be in common among Arabica, Robusta and
300 blends: 79/110; 97/110; 98/110; these fragments are characteristic, among others, of
301 pyridine, 5-methyl furfural, 1-methyl pyrrole, furfuryl pyrrole and furfuryl alcohol.
302 Each fragment varied linearly with color and, as a consequence, with roasting degree.
303 Table 3 gives correlation equations and coefficients, between indices and color, for
304 each variety/blend, together with the average index values and related standard
305 deviation. These index values can successfully be used to discriminate the degree of
306 roasting with good confidence, as is shown by their standard deviations that, within
307 the same variety, avoid any risk of two index intervals deriving from two different
308 roasting treatments (e.g. medium or dark) overlapping.

309 These results are particularly interesting for industrial applications, where in general
310 the material being processed comprises blends of green coffees with similar or

311 unvaried characteristics (variety, origins etc.); they enable a physical parameter
312 (color) to be correlated to chemical markers (indices) for the purpose of monitoring
313 the roasting process. Although the number of samples and varieties tested in the
314 present study is too small to be representative, these results outline the possibility of
315 defining indices of roasting and significant values for on-line process monitoring.
316 In addition, robust and reliable mathematical models to predict color directly from the
317 total spectral fingerprint and/or index values can successfully be applied on-line, not
318 only to predict color, but also to connect it to the aroma composition. A PLS model
319 equation was built and internally cross-validated with the 55 samples of the training
320 set and verified with the 25 samples of the test set (table 4). The results show a close
321 correlation between indices (i.e. m/z ratios) and color, with 17 of 25 samples having a
322 residual color measure below ± 3 , and only two of them (C21 and C24) presenting a
323 difference between measured and predicted color of approximately -9 . Table 4
324 reports the PLS results, together with the parameters of the model equation, and
325 shows a close correlation within the training set (rval: 0.9399) and a very satisfactory
326 standard error in color validation (SEV: 2.68). When the model equation was applied
327 to the samples of the test set, as expected the correlation coefficient in prediction
328 decreased (rpred: 0.8416) and the standard error in prediction increased (SEP: 4.29),
329 although the values were still satisfactory. These results show a reliable correlation,
330 although a less uniform training set would be necessary to include a wider range of
331 variables (variety, origin, roasting conditions) in a single equation. The results are
332 very satisfactory if the diversity represented is considered, because the samples of the
333 training set were very different and their number was relatively small.
334 In conclusion, the results show that HS-SPME-MS for on/in-line control of coffee
335 roasting process is a promising approach, through which not only the evolution of the
336 total MS profile can be studied, but also specific ions or ion ratios. The quick non-
337 separative method (HS-SPME-MS) described shows that a correlation between
338 spectral fingerprinting or roasting indices and color can be found, and that chemical
339 parameters can be used reliably to evaluate the degree of roasting. The combination
340 of MS profile and chemical indices with physical parameters affords more reliable
341 off-line monitoring or optimization of the roasting process, in particular to control
342 aroma development when developing new blends, and to detect the formation of toxic
343 compounds. In addition, the reliability of these results may be exploited as a
344 reference to validate those obtained by coupling a laboratory roasting machine to a
345 mass spectrometer directly, for on-line monitoring of the roasting process and marker
346 development³⁵⁻³⁸ HS-SPME-MS can also be used as analytical decision maker, i.e. to
347 select those sample(s) that have to be analyzed by conventional separative methods,
348 for instance when the non-separative intensities of some m/z fragments, diagnostic of
349 certain components, are outside the fixed limits of acceptance, or when the aroma
350 profile is not in line with that of the desired final product. Further studies are now
351 under way to extend the applicability of this method from model experiments to real-
352 world samples.

353

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358 indebted.

359

360 **SUPPORTING INFORMATION AVAILABLE**

361 One-way Anova results on fiber performance evaluation of the total volatile fraction,
362 spectral fingerprint areas of coffee pod samples, and method precision over 12
363 months, are included as supporting material. This material is available free of charge
364 via the internet at <http://pubs.acs.org>.

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Figure captions

Figure 1 HS-SPME-GC-MS (a), HS-SPME-TIC-MS (b) profiles and mass spectral fingerprint (c) of an Arabica coffee sample.

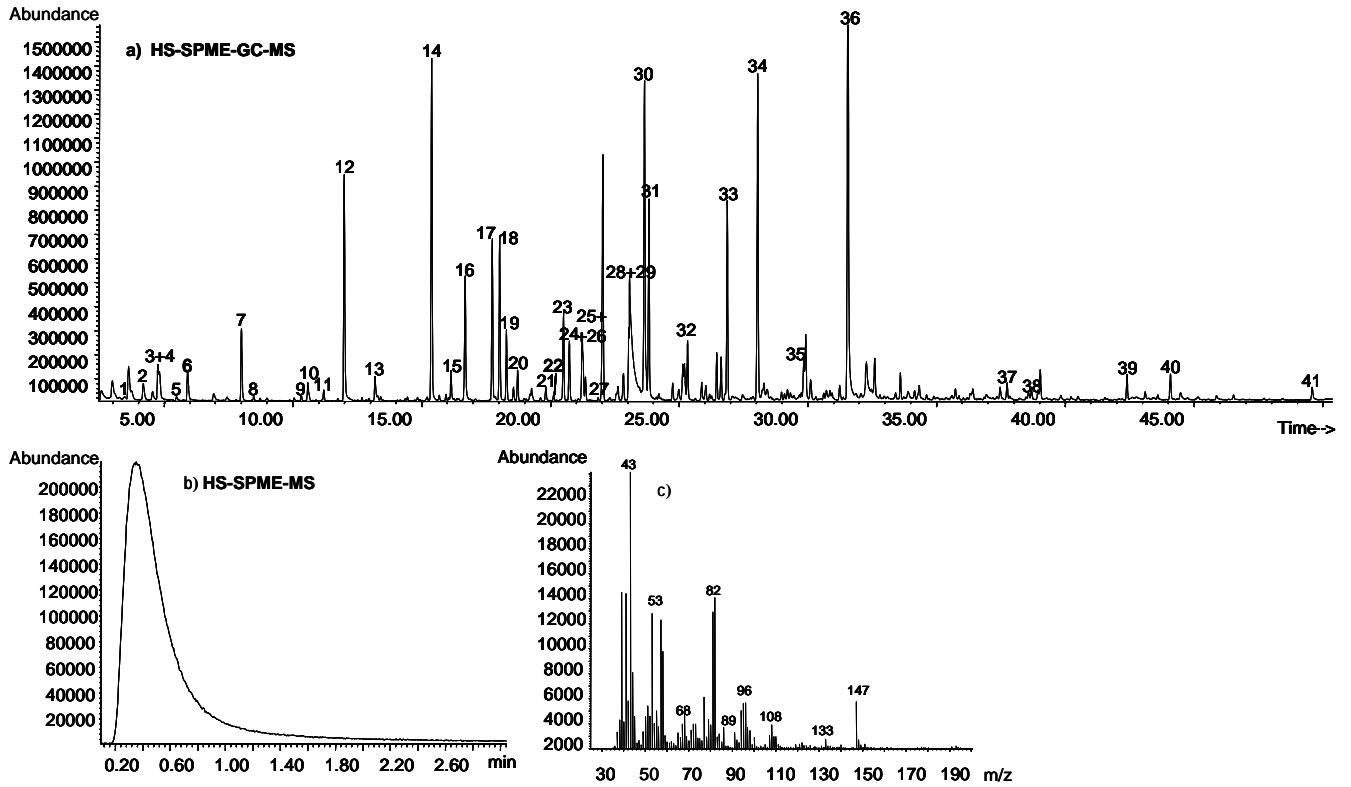
Figure 2 Scores (a) and Loading plots (b) of 55 different roasted coffee fingerprints (first 3PCs exp. var. 94.36%). Pre-processing: mean-center, full-cross validation. Categories: light roasting (solid triangle; color 57÷62), medium roasting (empty diamond; color 46÷53), dark roasting (solid diamond; colour 35÷42). Robusta: solid line; blend 50/50: dotted line; Arabica: dashed line.

Figure 3 Regression model for furan and 2-methylfuran on 55 coffee samples *versus* the degree of roasting, here represented by the experimentally-measured color.

Figure 4 OPLS regression model for color prediction as an association measure between volatile fraction, spectral fingerprint, and color of 25 commercial samples, whose origins, varieties and blends are unknown. ^a Correlation coefficient in Prediction, ^b Standard error in Prediction, ^c Correlation coefficient in Validation, ^d Standard error in Validation

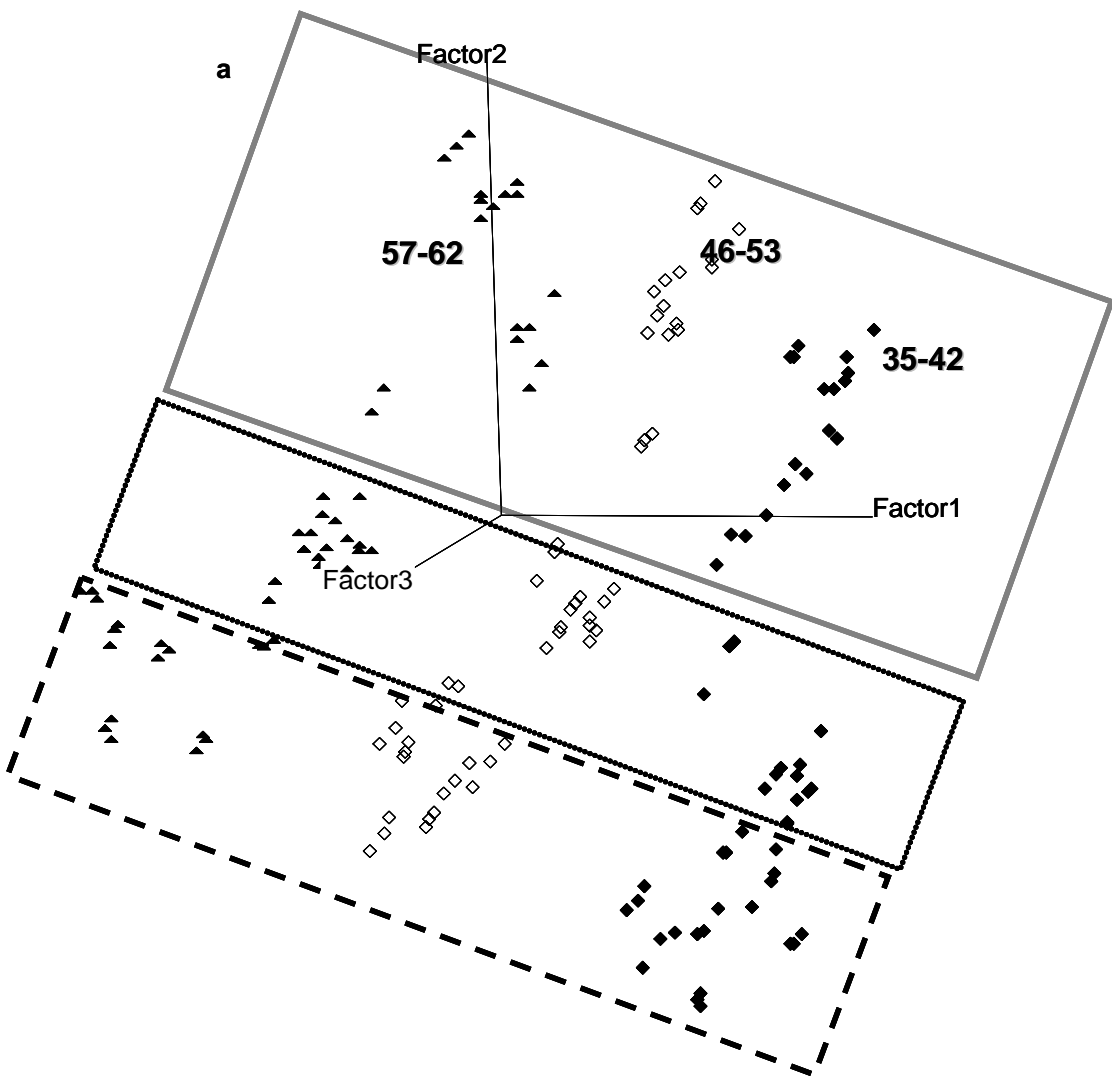
473 Figure 1 HS-SPME-GC-MS (a), HS-SPME-TIC-MS (b) profiles and mass spectral fingerprint (c) of an Arabica
474 coffee sample.

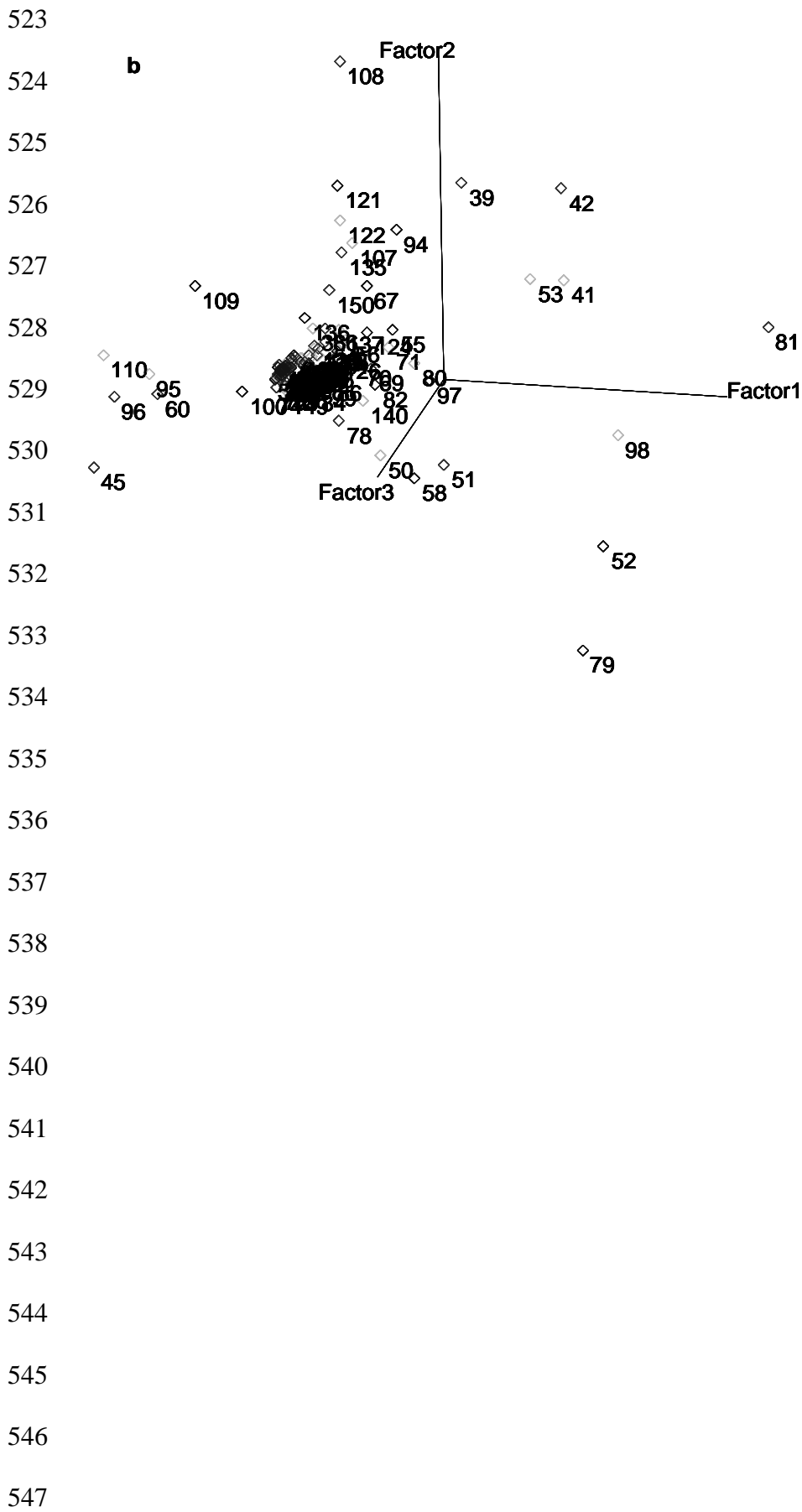
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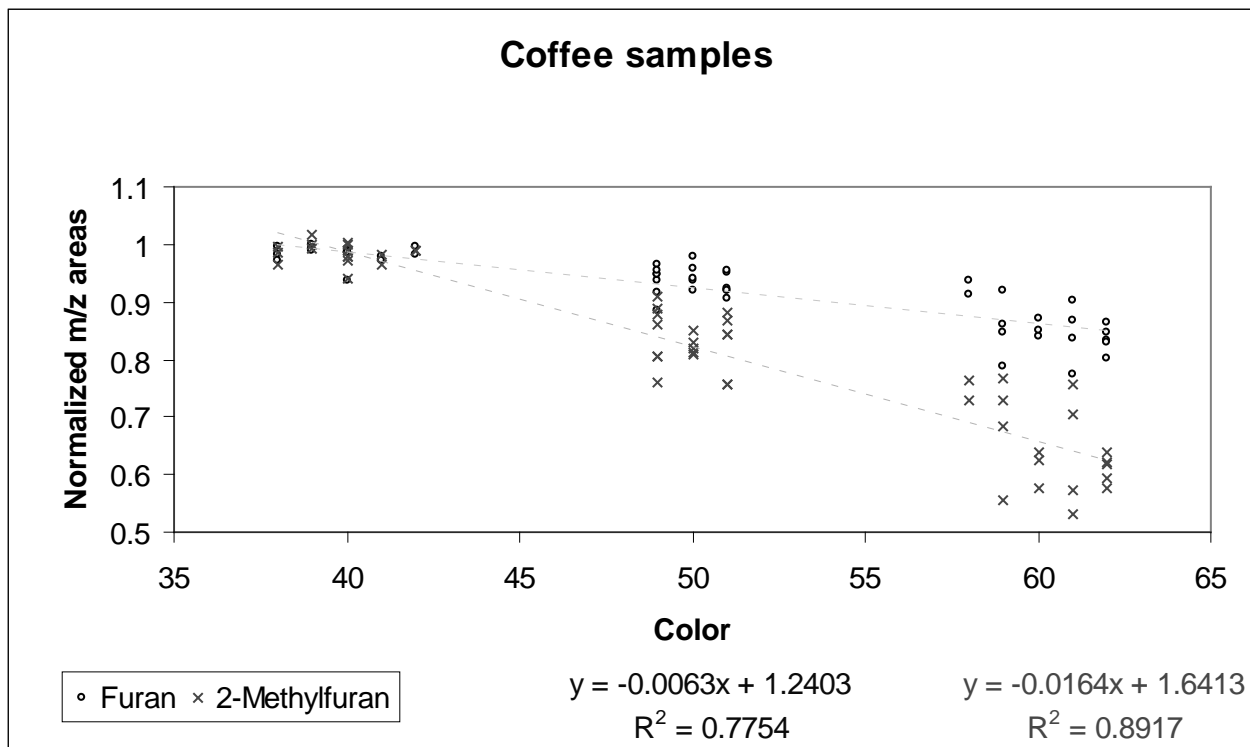
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548 Figure 3 Regression model for furan and 2-methylfuran on 55 coffee samples *versus* the degree of roasting
549 here represented by the experimentally measured color.

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564 Table 1 List of samples together with varieties and blends, color values and degrees of roasting .

Sample Code	Varieties and blends	Color Values	Roasting degree	Sample Code	Varieties and blends	Color Values	Roasting degree
Ara1	Arabica	61	Light	Rob13	Robusta	58	Light
Ara2	Arabica	51	Medium	Rob14	Robusta	50	Medium
Ara3	Arabica	40	Dark	Rob15	Robusta	40	Dark
Rob1	Robusta	61	Light	Ble13	Blend 50/50	60	Light
Rob2	Robusta	50	Medium	Ble14	Blend 50/50	49	Medium
Rob3	Robusta	40	Dark	Ble15	Blend 50/50	38	Dark
Ble1	Blend 50/50	59	Light	Ara16	Arabica	61	Light
Ble2	Blend 50/50	49	Medium	Ara17	Arabica	51	Medium
Ble3	Blend 50/50	39	Dark	Ara18	Arabica	42	Dark
Ara4	Arabica	59	Light	Rob16	Robusta	62	Light
Ara5	Arabica	49	Medium	Rob17	Robusta	50	Medium
Ara6	Arabica	40	Dark	Rob18	Robusta	40	Dark
Rob4	Robusta	60	Light	Ble16	Blend 50/50	62	Light
Rob5	Robusta	50	Medium	Ble17	Blend 50/50	49	Medium
Rob6	Robusta	41	Dark	Ble18	Blend 50/50	39	Dark
Ble4	Blend 50/50	61	Light	C1	Commercial	57	Light
Ble5	Blend 50/50	51	Medium	C2	Commercial	53	Medium
Ble6	Blend 50/50	42	Dark	C3	Commercial	47	Medium
Ara7	Arabica	62	Light	C4	Commercial	53	Medium
Ara8	Arabica	51	Medium	C5	Commercial	58	Light
Ara9	Arabica	39	Dark	C6	Commercial	39	Dark
Rob7	Robusta	62	Light	C7	Commercial	56	Light
Rob8	Robusta	49	Medium	C8	Commercial	58	Light
Rob9	Robusta	40	Dark	C9	Commercial	43	Dark
Ble7	Blend 50/50	60	Light	C10	Commercial	48	Medium
Ble8	Blend 50/50	51	Medium	C11	Commercial	39	Dark
Ble9	Blend 50/50	40	Dark	C12	Commercial	52	Medium
Ara10	Arabica	62	Light	C13	Commercial	50	Medium
Ara11	Arabica	50	Medium	C14	Commercial	43	Dark
Ara12	Arabica	41	Dark	C15	Commercial	48	Medium
Rob10	Robusta	59	Light	C16	Commercial	61	Light
Rob11	Robusta	51	Medium	C17	Commercial	52	Medium
Rob12	Robusta	38	Dark	C18	Commercial	47	Medium
Ble10	Blend 50/50	58	Light	C19	Commercial	56	Light
Ble11	Blend 50/50	49	Medium	C20	Commercial	40	Dark
Ble12	Blend 50/50	38	Dark	C21	Commercial	47	Medium
Ara13	Arabica	59	Light	C22	Commercial	57	Light
Ara14	Arabica	49	Medium	C23	Commercial	48	Medium
Ara15	Arabica	40	Dark	C24	Commercial	46	Medium
				C25	Commercial	52	Medium

569 Table 2 List of markers identified in HS-SPME-GC-MS profile of an Arabica roasted sample together with their
 570 target (TI) and qualifier (Qi) ions. *Markers tentatively identified through their MS-EI fragmentation patterns

ID	Compounds	Ret Time (min)	I_{CW}^T	I_{OV1}^T	TI	Qi1	Qi2
1	Furan	3.74	837	500	68	58	39
2	2-methylfuran	4.49	873	586	82	81	53
3	2-methyl butanal	5.09	903	641	86	57	41
4	3-methyl butanal	5.09	904	635	86	71	57
5	2,5-dimethylfuran	5.86	938	691	96	95	81
6	2,3-butanedione	6.31	960	555	86	57	43
7	2,3-pentandione	8.49	1043	668	100	57	43
8	2-vinylfuran*	9.00	1059	-	94	65	66
9	2,3-hexanedione	10.88	1117	756	43	71	43
10	1-methyl-pyrrole	11.18	1124	715	81	80	66
11	2-vinyl-5-methylfuran*	11.79	1139	-	108	107	79
12	Pyridine	12.62	1165	720	79	52	39
13	Pyrazine	13.85	1195	709	80	53	70
14	methylpyrazine	16.08	1249	802	94	67	53
15	3-hydroxy-2-butanone	16.84	1265	681	88	73	45
16	1-hydroxy-2-propanone	17.42	1278	626	74	43	41
17	2,5-dimethylpyrazine	18.45	1306	893	108	81	42
18	2,6-dimethylpyrazine	18.74	1313	889	108	81	42
19	ethylpyrazine	19.01	1318	895	107	108	80
20	2,3-dimethylpyrazine	19.45	1329	904	108	67	93
21	1-hydroxy-2-butanone	20.57	1353	732	88	57	42
22	3-ethyl-pyridine	20.85	1364	934	107	92	79
23	2-ethyl-6-methylpyrazine	21.23	1371	977	121	122	94
24	2-ethyl-5-methylpyrazine	21.46	1376	981	121	122	94
25	trimethylpyrazine	21.96	1389	984	122	81	42
26	2-ethyl-3-methylpyrazine	22.01	1391	985	121	122	80
27	2-propylpyrazine	22.60	1402	985	94	107	122
28	2-ethyl-3,6-dimethyl pyrazine	23.78	1427	1059	135	136	108
29	acetic acid	23.79	1432	547	60	43	45
30	furfural	24.45	1443	801	96	95	39
31	1-acetoxy-2-propanone	24.57	1448	825	43	86	116
32	2-acetylfuran	26.12	1483	882	95	110	39
33	furfuryl acetate	27.64	1521	963	81	98	140
34	5-methyl furfural	28.86	1551	933	110	109	81
35	1-methyl-2-carboxaldehyde pyrrole	30.67	1596	974	109	108	80
36	furfuryl alcohol	32.37	1640	823	98	81	69
37	1-furfuryl pyrrole	38.58	1805	1152	81	147	53
38	guaiacol	39.54	1832	1064	109	124	81
39	2-acetyl pyrrole	43.27	1941	1030	94	109	66
40	2-carboxaldehyde pyrrole	44.98	1991	976	95	94	66
41	p-vinylguaiacol	50.47	2163	1289	150	135	107

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573 Table S1 One-way ANOVA results on Total Volatile fraction spectral fingerprint areas of coffee pod samples
 574 (n=5; $\alpha=0.05$).

<i>Groups</i>	<i>Count</i>	<i>Sum</i>	<i>Mean</i>	<i>Variance</i>
Fibre 1 (F1)	5	103.41	20.68	2.38
Fibre 2 (F2)	5	134.87	26.97	0.22
Fibre 3 (F3)	5	103.14	20.63	0.52

One-Way Anova

<i>Source of variation</i>	<i>SQ</i>	<i>gdl</i>	<i>MQ</i>	<i>F</i>	<i>Significant Value</i>	<i>F crit</i>
Between groups	133.08	2	66.54	63.86	4.01E-07	3.89
Within groups	12.50	12	1.04			
Total	145.59	14				

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577 Table S2 Method precision (repeatability and intermediate precision) of total fingerprint and some
 578 characteristic ions expressed as Relative Standard Deviation % calculated over replicates collected across the
 579 entire period (12 months). * m/z ions characterizing key-aroma compounds according to Czerny²³ and Blank²⁴.
 580

<i>m/z</i>	<i>Characterizing properties</i>	<i>Repeatability</i>	<i>Intermediate precision</i>
37	Roasting	1.30	14.15
38	Roasting	1.10	9.32
39	Aroma and roasting	0.38	5.06
41	Aroma and roasting	0.84	16.27
45	Aroma and varieties	2.25	4.65
50	Aroma and roasting	0.36	9.29
51	Aroma and roasting	1.06	8.04
52	Aroma and roasting	1.72	8.50
53	Aroma and roasting	2.27	6.50
60	Aroma and varieties	1.70	8.00
62	Roasting	1.63	8.00
66	Roasting and varieties	1.95	9.28
67	Roasting	0.99	8.48
68	Toxic	1.92	7.84
69	Aroma and roasting	1.70	10.37
70	Aroma and roasting	2.19	22.77
78	Roasting	1.67	9.29
79	Roasting	1.22	12.67
80	Roasting	0.58	10.35
81	Roasting and varieties	1.60	9.10
82	Toxic	0.58	6.50
86*	Aroma	2.62	7.62
88	Aroma and varieties	4.62	16.12
94	Aroma and roasting	1.14	13.56
95	Roasting and varieties	1.29	7.68
96	Roasting	2.22	9.82
97	Roasting	2.27	5.26
98	Roasting	2.31	7.67
100*	Aroma	4.47	13.74
108	Aroma and roasting	0.50	11.96
109*	Aroma and roasting	1.34	6.98
110	Roasting	1.70	7.86
121	Aroma and roasting	2.49	9.98
122	Aroma and roasting	2.54	11.32
135*	Aroma	2.44	5.19
150*	Aroma	11.24	20.08
Total Fingerprint Area		1.27	3.16

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583 Table 3 Indices (m/z ratios) and related correlation equation and coefficients with the color for each variety
 584 and/or blend analyzed together with each index value and its standard deviation.

585

Index m/z ratios	Variety or blend	Equation	r ²	Index value	Index Standard deviation	Mean color value
79/110	100% ARABICA	$y = -119x + 8018$	0.8754	1051	226	60
				1765	358	50
				3426	600	40
	BLEND	$y = -110x + 7604$	0.9872	1161	207	60
				1796	310	50
				3361	343	40
	100% ROBUSTA	$y = -63x + 4874$	0.9550	1148	111	60
				1621	135	50
				2407	123	40
97/110	100% ARABICA	$y = -51x + 3884$	0.8946	879	76	60
				1239	143	50
				1898	139	40
	BLEND	$y = -48x + 3872$	0.9912	1013	101	60
				1382	154	50
				1979	139	40
	100% ROBUSTA	$y = -41x + 3571$	0.9925	1148	118	60
				1529	90	50
				1958	107	40
98/110	100% ARABICA	$y = -108x + 8043$	0.8896	1692	172	60
				2438	303	50
				3846	325	40
	BLEND	$y = -101x + 7913$	0.9918	1960	206	60
				2725	333	50
				3971	285	40
	100% ROBUSTA	$y = -81x + 7034$	0.9939	2203	252	60
				2967	171	50
				3818	229	40

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588 Table 4 Color measured with the three indices through PLS elaboration and model parameters

sample	Measured Y	Predicted Y	Residual Y	Upper Limit	Lower Limit
Ara2	51	52	-0.71	57	46
Rob3	40	45	-4.59	50	39
Ble1	59	56	2.73	61	51
Ara5	49	49	0.02	54	44
Ble4	61	56	4.82	61	51
Rob7	62	57	5.13	62	52
Rob8	49	48	1.12	53	43
Ara10	62	59	2.74	65	54
Rob10	59	54	5.18	59	49
Ble12	38	38	-0.43	44	33
Ara15	40	40	0.32	45	34
Ble13	60	57	3.21	62	52
Ble14	49	48	0.80	53	43
Ara18	42	42	0.08	47	37
Rob18	40	43	-2.90	48	38
Ble16	62	58	4.25	63	53
C19	56	62	-5.84	67	57
C21	47	57	-9.79	62	52
C24	46	55	-9.26	60	50
C14	43	36	6.66	42	31
C17	52	52	-0.43	58	47
C1	57	56	1.27	61	50
C4	53	54	-0.91	59	49
C6	39	41	-2.05	47	36
C10	48	55	-6.76	60	50
C12	52	53	-1.22	58	48
Prediction Equation y=0.754x+12.71	rpred 0.8416	SEP 4.29	Model validation error on 2 Factors	rval 0.9399	SEV 2.68

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