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# CHARACTERIZATION OF CRUMB RUBBER FROM END-OF-LIFE TYRES FOR PAVING APPLICATIONS

M.C. ZANETTI, S. FIORE, B. RUFFINO, E. SANTAGATA, D. DALMAZZO, M. LANOTTE

*Department of Environment, Land and Infrastructure Engineering, Politecnico di Torino – Corso  
Duca degli Abruzzi 24, 10129, Torino, Italy.*

*E-mail: [mariachiara.zanetti@polito.it](mailto:mariachiara.zanetti@polito.it). Tel. 00390115647696.*

**SUMMARY:** Crumb rubber (CR) derived from grinding of end-of-life tyres (ELTs) may be successfully used as a bitumen modifier or as a supplementary component in the production of bituminous mixtures employed for the construction and maintenance of road pavements. However, CRs deriving from different sources and processes yield effects on performance under traffic loading and on gaseous emissions produced during laying on site which may change considerably depending upon their physical and chemical properties. In order to quantitatively assess the possible variability of CR characteristics, samples were taken from 9 Italian and 2 foreign ELT processing plants. Investigation activities included field surveys, during which plants were examined in detail, and laboratory tests, which focused on physical and chemical characterization of CR. Based on the analysis of available technical information and experimental data, it was possible to find relationships between the peculiar characteristics of treatment cycles and corresponding CR properties.

**KEY WORDS:** crumb rubber, end-of -life tyres, paving applications, physical chemical characterization

## 1. INTRODUCTION

27 Management of end-of-life tyres (ELTs) has become a critical problem worldwide due to the  
28 increasing number of vehicles circulating in the road network and to the crucial role that mobility has  
29 assumed in society development. Since landfill disposal has been banned in most Countries,  
30 alternative final destinations have been sought, with a major effort being placed in trying to exploit  
31 in the most efficient manner the high energy potential of ELTs. Nevertheless, due to the fact that  
32 rubber employed in tyre fabrication is the result of specialized materials' selection, recycling and  
33 reuse seem to be preferable options for such a high-quality waste material (Santagata and Zanetti,  
34 2012).

35 Practical experience and research have shown that crumb rubber (CR) derived from grinding of  
36 ELTs may be successfully used as a bitumen modifier or as a supplementary component in the  
37 production of bituminous mixtures employed for the construction and maintenance of road  
38 pavements. Available technologies can be grouped into two main categories which are associated to  
39 the so-called "wet" and "dry" production processes. Within each group, different versions of the  
40 technology have been conceived and subjected to trials either in the laboratory or at the industrial  
41 scale, in the constant attempt of exploiting more efficiently the performance-related benefits of CR.

42 In the "wet" process, CR is preliminarily mixed with bitumen, thus obtaining a ductile and elastic  
43 modified binder, known as "asphalt rubber" (ASTM D6114-09), that is then combined with  
44 aggregates in the hot mix plant. Resulting mixtures are generally of the gap-graded (GG) or open-  
45 graded (OG) type, characterized by a very high binder content (of the order of 7.5-10% b.w. of dry  
46 aggregates) and by a non-continuous particle size distribution that allows CR to be accommodated  
47 within the composite material. GG and OG mixtures are employed for the formation of surface  
48 courses and have earned a satisfactory reputation with respect to field performance (Hicks, 2002).

49 In the "dry" method, CR is introduced in the production flow of bituminous mixtures as a  
50 supplementary component, substituting part of the aggregates and providing enhanced elastic  
51 response under loading (Santagata and Zanetti, 2012; Santagata et al., 2013). Mixtures are usually of  
52 the dense-graded (DG) type, with a continuous particle size distribution and an optimal binder

53 content (usually of the order of 5-6%) which is only slightly higher than that adopted for standard  
54 mixtures containing no recycled rubber (Buncher, 1995). Unfortunately, the performance record of  
55 these mixtures has been quite inconsistent, with the frequent occurrence of early ravelling  
56 phenomena and moisture-related damage (Amirkhanian, 2001; Caltrans, 2005). This also explains  
57 the limited diffusion of such a technology, with full-scale applications that have been generally  
58 carried out locally rather than at the network level.

59 For both the abovementioned technologies, concerns have been raised on the use of CR in  
60 bituminous mixtures with respect to its potential contribution to gaseous emissions during production  
61 and laying, and to the possible consequences which it can cause on the health of construction  
62 workers. However, a limited number of experimental studies have been carried out on this specific  
63 topic, with no clear quantification of the actual hazardous effects of CR (Watts et al., 1998; Burr et  
64 al., 2001; Stout and Carlson, 2003).

65 The Authors have recently contributed to this area of technical knowledge by applying to pavement  
66 works risk analysis concepts developed in previous studies focused on the evaluation and  
67 remediation of contaminated sites (Marescalco and Zanetti, 2010; Zanetti et al., 2013b) and on the  
68 approval of the use of CR in artificial turf sports fields (Ruffino et al., 2013). In particular, by  
69 considering the results of analyses carried out on gaseous emissions sampled on site or in the  
70 laboratory in controlled conditions, comparisons have been made between bituminous mixtures  
71 containing CR (produced by employing both the “wet” and “dry” technology) and bituminous  
72 mixtures of the standard type (Zanetti et al., 2013a, 2014a and 2014b).

73 Based on the Authors’ experience, it can be concluded that CRs deriving from different sources and  
74 processes yield effects on field performance and emissions of bituminous binders and mixtures  
75 which may change considerably depending upon their physical and chemical properties. In turn,  
76 these are dictated by the characteristics of ELTs subjected to processing and by the phenomena  
77 which occur during the various phases of ELT treatment (shredding, magnetic separation,  
78 granulation, milling and sieving).

In order to quantitatively assess the possible variability of CRs, in the study described in this paper, samples were taken from 9 Italian and 2 foreign ELT processing plants and thereafter subjected to laboratory investigations for the assessment of physical and chemical characteristics. In particular, experimental tests were carried out for the determination of particle size distribution, density, cleanliness, particle shape and morphology, specific surface area, content of metals, PAHs (polynuclear aromatic hydrocarbons) and VOCs (volatile organic compounds) and elemental analysis. Based on the analysis of technical information and experimental data, it was possible to find relationships between the peculiar characteristics of treatment cycles and corresponding CR properties.

## **2. BACKGROUND**

### **2.1 CR production**

Processing of ELTs is carried out in various phases during which rubber is separated from other materials (textile fibers and metals), mechanically reduced in granular form and finally divided into particle size fractions. When shredding, shear and abrasion operations occur with no specific temperature conditioning, the process is known as “ambient size reduction”. In some plants, however, ELTs are brought below rubber glass transition temperature with adequate cooling systems and the resulting process is referred to as “cryogenic”. Moreover, novel processing methods are continuously developed in order to optimize plant efficiency. As an example, size reduction has also been attempted by means of the “high pressure waterjet” system, based on the abrasive effects caused by water jets at 3.000 bar which pulverize ELTs.

Depending upon the type of treatment process and on the origin of ELTs fed to it, CR may have different physical and chemical characteristics. In particular, particles deriving from ambient size reduction generally have irregular shape and rough surface; moreover, it has been postulated that in some cases heat generated during mechanical processing may induce a partial devulcanization of rubber. By comparison, cryogenically produced CRs are mostly made of cuboid-shaped particles

105 with a smooth surface. With respect to ELT origin, it is well known that tyre producers employ  
106 different rubber formulations and that truck tyres generally have a higher natural rubber content than  
107 car tyres. However, in practice only the second factor may influence CR production since there are  
108 plants that treat only truck (or car) tyres, but none are dedicated to a single tyre producer.

## 109 **2.2 CR-bitumen interaction**

110 In the asphalt rubber “wet” production process, CR is thoroughly mixed with bitumen at a  
111 temperature in the 175-225°C range. The resulting binder is then kept in agitation at high  
112 temperature (150-215°C) for the time period (of the order of 45-60 minutes) which is necessary for  
113 interaction phenomena between the components to occur. In particular, CR particles are partially  
114 digested in the bituminous matrix and absorb part of the aromatic fraction of bitumen, with a  
115 resulting volume expansion and formation of a gel-like surface coating which gives the binder its  
116 peculiar physical and rheological characteristics (Way et al., 2012). In this form, CR particles are  
117 still visible in the composite binder which has a distinctive granular-like appearance. If curing is  
118 carried out at an excessive temperature and/or for a too long time period, degradation phenomena  
119 become prevalent and CR is totally digested in bitumen: as a consequence, the resulting binder does  
120 not have the typical characteristics of asphalt rubber and may exhibit unsatisfactory performance.  
121 Asphalt rubber binders usually have a CR content comprised between 18 and 22% (b.w. of total  
122 binder), with a high viscosity at storage/mixing temperatures and enhanced elastic properties in  
123 service. Binder characteristics are dependent not only upon chemical composition of employed  
124 components, but also on CR dosage, particle size and morphology. In such a context, it has been  
125 proven that the intensity of the above described interaction phenomena tends to increase with CR  
126 dosage and specific surface area (Shen et al., 2009). Therefore, CRs which are considered more  
127 reactive are those which are finer, constituted by rough, irregular particles.

128 In the production of “dry” mixtures CR is usually employed with a dosage comprised between 1%  
129 and 3% (b.w. of dry aggregates). Depending upon the type of plant, either batch or drum-mix, CR  
130 can be introduced in the production flow of bituminous mixtures by means of different methods, but

is always added to the heated aggregates before coming in contact with bitumen. When this condition occurs, even though CR particles are not digested in bitumen, they do absorb part of its aromatic fractions. However, such an interaction takes place in non-controlled conditions, starting from the mixing process in the plant and progressing throughout the early phases of service life (Santagata et al., 2013). Studies performed in the past have shown that in this respect beneficial effects can be obtained by pretreating CR by means of function-specific catalysts (Epps, 1994) or extender oils (Newcomb et al., 1994; Khalid and Artamendi, 2002).

### **2.3 CR selection and acceptance**

Selection and acceptance of CR for use in asphalt rubber classically relies upon the requirements that have been set in ASTM D6114. These are expressed in terms of cleanliness (fiber content < 0.5%; metal content < 0.01%), moisture content (< 0.75%), density (equal to  $1.15 \pm 0.05$ ) and maximum particle size (2.36 mm). However, the standard also specifies that the exact size distribution of CR should be agreed upon between producer and end-user.

Based on experience and local tradition, in other contexts requirements may be slightly different. As an example, technical specifications of the California State Department of Transportation require the use of natural and synthetic rubber in given proportions and provide size distribution acceptance intervals which should be satisfied by single CR components (Caltrans, 2003).

In the case of the “dry” production technology, depending upon the desired effect on the resulting bituminous mixture, CR can be employed with different size distributions, ranging from “ultrafine” (entirely passing the 0.3 mm sieve) to “coarse” (passing the 6 mm sieve and totally retained on the 2.36 mm sieve) (Caltrans, 2005). Since it has been recognized that size distribution (and related specific surface area) is the main factor controlling performance properties of “dry” mixtures, specific requirements are usually not set on other CR characteristics.

Physical and rheological properties of asphalt rubber are also subjected to acceptance requirements in technical specifications. This is done either by referring to the results of classical empirical tests (penetration, softening point, etc.) or by considering viscoelastic properties measured at

representative temperatures, frequencies and ageing conditions. All evaluation systems also include acceptance criteria referred to apparent viscosity, which is typically required to be comprised between 1500 and 5000 mPa·s. Such a property, which refers to the flow behavior of asphalt rubber, is considered as a good quality indicator that can be also measured on site, from storage tanks, with portable hand-held viscometers.

### **3. EXPERIMENTAL INVESTIGATION**

The investigation described in this paper was carried out by considering 16 CRs sampled from 9 Italian and 2 foreign plants. Experimental activities included field surveys, during which processing plants were examined in detail, and laboratory investigations, which focused on CR physical and chemical characterization. Tests were carried out in the “Road Materials Laboratory” (RML) and in the “Environmental Chemistry Laboratory” (ECL) of the Politecnico di Torino

#### **3.1 Plant surveys**

Treatment processes of the plants which were considered in this study are synthetically described in Table 1. The majority of them (9 out of 11) operate by mechanical size reduction at ambient temperature, one is of the cryogenic type (G) and one relies upon the waterjet technology (H). With respect to ambient size reduction processes, it can be observed that the number and type of consecutive working phases changes considerably, thus affecting the characteristics of final products. In general terms, more recent plants (e.g. B and E) tend to have multiple iron magnetic separation phases in order to ensure a greater purity of CRs; moreover, multiple shredding/milling phases may be combined with the purpose of increasing production flexibility (i.e. of widening the range of possible CR products).



183 Table 1. Description of ELT processing plants.

| Phases                   | ELT processing plants |   |   |   |   |   |   |   |      |   |   |
|--------------------------|-----------------------|---|---|---|---|---|---|---|------|---|---|
|                          | A                     | B | C | D | E | F | G | H | I    | J | K |
| Primary shredding        | ×                     | × | × | × | × | × | × | × | n.a. | × | × |
| Iron magnetic separation |                       |   |   |   | × |   |   |   | n.a. | × |   |
| Secondary shredding      |                       | × |   |   | × | × | × |   | n.a. |   | × |
| Cold granulation         |                       | × | × |   |   |   |   |   | n.a. | × |   |
| Iron magnetic separation | ×                     | × | × | × | × | × | × |   | n.a. |   | × |
| Primary milling          | ×                     | × | × | × | × | × | × |   | n.a. | × |   |
| Secondary milling        | ×                     |   |   |   |   |   |   |   | n.a. |   |   |
| Sieving                  | ×                     | × | × | × | × | × | × | × | n.a. | × | × |

(\*) Carried out in cryogenic conditions

(\*\*) Carried out with waterjet treatment.

n.a. Information not available.

### 184 3.2 CR chemical characterization

185 Chemical characterization of CR consisted in the determination of the contents of metals, volatile  
186 organic compounds (VOCs), polycyclic aromatic hydrocarbons (PAHs) and elemental analysis  
187 (carbon, hydrogen, nitrogen and sulfur). Since there are no fixed standards which define procedures  
188 for CR characterization, tests were carried out by following protocols which were developed for such  
189 a purpose in the ECL. In all cases, duplicate measurements were performed for each test.

190 Metals were determined by using an inductively coupled plasma optical emission spectrometer (ICP-  
191 OES). Test samples were obtained by subjecting 0.1 g of CR to treatment in a microwave digestion  
192 unit in the presence of 1 ml perchloric acid (70%, Merck) and 3 ml of nitric acid (65%, Riedel de  
193 Haen, Sigma Aldrich). After digestion, samples were filtered (Whatman filters, 2.7 micron retention  
194 grade) and then transferred to 100 ml flasks where they were brought to full volume by means of  
195 distilled water. In ICP-OES analyses, elements which were detected through their electromagnetic  
196 radiation after excitation included those which derive from residues of the metal structure of tyres  
197 and those which are present due to contamination.

198 Determination of VOCs and PAHs was carried out by means of solvent extraction and  
199 gas-chromatographic analysis. A 2 g CR sample was extracted with 20 ml of CH<sub>2</sub>Cl<sub>2</sub> kept for 20  
200 minutes in a microwave oven set at 600 W. Analyses were then performed by using an Agilent  
201 7890/5975 gas chromatograph equipped with a HP5-MS capillary column (30m×0.25mm×0.25μm)  
202 and combined with a mass spectrometer detector (GC-MS). VOC and PAH compounds were  
203 considered in the investigation since they are potentially toxic or carcinogenic substances.  
204 Carbon, hydrogen, nitrogen and sulphur contents were determined by employing a Flash 2000  
205 ThermoFisher Scientific CHNS analyzer which operates according to the dynamic flash combustion  
206 technique on 2-3 mg CR samples. After combustion of the sample, reaction gas products were  
207 carried by helium flow to a copper-filled layer, then through a GC column that provided separation  
208 of the combustion gases and finally detected by a Thermal Conductivity Detector.

### 209 **3.3 CR physical characterization**

210 Physical characterization of CR was carried out by means of laboratory tests for the determination of  
211 particle size distribution, density, cleanliness, particle morphology and specific surface area.  
212 Particle size distribution was evaluated in dry conditions by making use of sieves of the Tyler series  
213 (ASTM E-11, 2000). Since electrostatic effects may make this type of analysis quite difficult,  
214 especially in the presence of high percentages of very fine material, intense mechanical agitation of  
215 the sieve column was required. At least three repetitions were considered necessary in order to have  
216 representative results. Data was also cross-checked with those derived from image analysis,  
217 described further on in the context of morphology evaluation.  
218 Density ( $\rho$ ) at 25°C was measured with the pycnometer method (EN1097-7, 2008) by employing  
219 ethylic alcohol as fluid of known density in order to prevent particles from floating to the surface.  
220 Relative density (i.e. specific gravity, SG) was thereafter calculated by referring measured density to  
221 that of water at the same temperature.

222 Evaluation of the degree of cleanliness (presence of textile fibers, metal residues and other  
 223 contaminants) was performed in qualitative terms by means of observations with a stereomicroscope.  
 224 For such a purpose, images were acquired with a digital camera and thereafter visually examined.  
 225 Use of stereomicroscope observations was also the starting point of the technique used for the  
 226 assessment of morphological characteristics and for the estimate of surface area (Santagata et al.,  
 227 2012). However, in this case the plan-view digital image of the set of considered particles was  
 228 processed with a freeware software (ImageJ, version 1.45, National Institutes of Health) which  
 229 allows the identification and geometrical description of each particle.

230 The following morphological parameters were directly calculated from image analysis results:

- 231 ▪ average value of the shape coefficient ( $C_f$ ), given by the ratio between the maximum and  
 232 minimum Feret diameters ( $\max F$  and  $\min F$ ) of each particle;
- 233 ▪ average value of the solidity coefficient ( $C_s$ ), given by the ratio between the area of each particle  
 234 ( $A_{\text{particle}}$ ) and the minimum convex area ( $A_{\text{convex}}$ ) in which it is enclosed.

235 Surface area per unit mass ( $SA_m$ ) was calculated by making use of the following expression:

$$236 \quad SA_m = \phi \cdot \frac{6}{\rho} \cdot \sum_i \frac{f_i}{d_{m,i}} \quad (1)$$

238 where:  $SA_m$  = surface area per unit mass (in  $m^2/g$ );

239  $\phi$  = corrective factor which takes into account morphology of constituent particles;

240  $\rho$  = density (in  $g/m^3$ );

241  $f_i$  = frequency (in decimal units) of the  $i$ -th size fraction;

242  $d_{m,i}$  = mean particle diameter (in m) of the  $i$ -th size fraction.

243 Corrective factor  $\phi$  was derived from image analysis results by making use of analytical models  
 244 which allow calculation of surface area per unit volume ( $SA_v$ ) by referring to sets of ideal particles  
 245 with different shape and roughness characteristics (Santagata et al., 2012). In particular,  $\phi$  was

246 obtained as the product of two distinct corrective factors,  $\phi_f$  and  $\phi_r$ , which separately take into  
247 account the effects of shape and roughness, respectively.

248

#### 249 **4. RESULTS AND DISCUSSION**

250 Results recorded during laboratory tests are synthesized in Tables 2-7 and in Figures 1-7. In general  
251 terms, as expected, it was found that sampled CRs represent a very broad spectrum of products for  
252 paving applications which may constitute a valuable reference database for future use. As discussed  
253 in the following, data analysis was carried out by highlighting relevant differences between products  
254 and by searching for relationships between investigated characteristics.

255 By taking into account measured size distribution, as required by the EN standard currently under  
256 development (CEN/TS 14243, 2010), CR samples were associated to alphanumeric codes given by  
257 the label of the production plant (from A to K) and the interval comprised between  $D_{10}$  and  $D_{90}$   
258 (sieve openings corresponding to 10% and 90% passing, respectively). Thus, it was found that  
259 considered CRs can be grouped into three categories: “coarse” ( $D_{90} > 1.0$ ), “standard” ( $0.5 \leq D_{90}$   
260  $1.0$ ) and “fine” ( $D_{90} < 0.5$ ). In such a context it should be mentioned that CR H 0-0.7, derived from  
261 waterjet production, although technically allocated to the “standard” category, has very peculiar  
262 characteristics as a consequence of its very broad size distribution which combines the presence of  
263 coarse particles (with size up to 2 mm) with a very high content of fines (14.3% passing the 0.063  
264 sieve).

265 Particle size distribution curves are displayed in Figures 1a and 1b, while Table 2 lists the values of  
266 descriptive size distribution parameters which include  $D_{10}$  and  $D_{90}$ ,  $D_{50}$  (which provides a measure of  
267 the overall coarseness/fineness),  $D_{80}/D_{20}$  (a uniformity factor which quantifies overall width of size  
268 distribution) and  $P_{0.063}$  (percent passing the 0.063 sieve, corresponding to the content of the very fine  
269 particles).

270 It can be observed that among the “standard” CRs, considered as typical candidates for use in the  
271 production of asphalt rubber (“wet” process), the main factor which allows them to be discriminated

272 is  $P_{0.063}$ , which in fact is especially high in the case of the waterjet (H 0-0.7) and cryogenic  
273 (G 0.2-0.6) products. This observation proves that within a given category of CRs, significant  
274 differences in terms of size distribution can generally stem only from variations of production type.  
275 Similar observations apply to the category of “fine” CRs, usually employed for special applications  
276 as a result of their higher market value. It can be observed that plant B, which in comparison to the  
277 others (A and C) has a greater number of shredding/granulation/milling phases, yields the product  
278 with the highest percentage of very fine particles ( $P_{0.063}$  equal to 59.9%).  
279 No specific comments can be made on the “coarse” CRs due to the similarity of the two considered  
280 products and plants.

281 Figure 1a. Particle size distribution of the CR samples (plants A-E).

282 Figure 1b. Particle size distribution of the CR samples (plants F-K).

283 Basic physical characterization of CRs was completed by performing laboratory tests for the  
284 evaluation of density  $\rho$  (and specific gravity, SG). Corresponding results, listed in Table 2, are  
285 comprised within a narrow variation range ( $\rho = 1.158\text{-}1.223\text{ g/cm}^3$ ;  $\text{SG} = 1.160\text{-}1.225$ ), and show  
286 that most of the CRs of the “standard” category meet requirements set by ASTM ( $\text{SG} = 1.15 \pm 0.05$ )  
287 for products to be used for the production of asphalt rubber. The most relevant deviation is exhibited  
288 by cryogenic CR G 0.2-0.6, which has a very high density value ( $1.223\text{ g/cm}^3$ ), probably as a result  
289 of the peculiar morphological properties of its particles (cuboid-shaped and very smooth).  
290 Since CR-bitumen interaction and properties of the resulting binder/matrix are strongly dependent  
291 upon particle morphology and specific surface area, these characteristics were assessed by means of  
292 adequate models applied to image analysis results derived from microscope observations. Examples  
293 of post-processing images of two CRs are shown in Figure 2, while results of the consequent  
294 analytical evaluation are given in Tables 3a and 3b and in Figures 3 and 4.

295 Morphological coefficients  $C_s$  and  $C_f$  were calculated from the geometry of the projection of CR  
296 particles on the horizontal plane. While  $C_f$  refers to the degree of elongation,  $C_s$  reveals the presence  
297 of surface irregularities which are responsible for a significant increase of total perimeter (and thus,  
298 in the three-dimensional situation, of surface area).

299 It was observed that low  $C_f$  values, typical of regularly-shaped particles which are those desired for  
300 the preparation of homogeneous blends/mixtures, may be accompanied, depending upon the specific  
301 features of each CR (and of the originating production process), by either high or low values of  $C_s$ .  
302 This is clearly shown in the shape-roughness mapping plot of Figure 3, built by representing  $C_s$  as a  
303 function of  $C_f$ . Most of the considered CRs fall within the cuboid-rough (low  $C_f$ , low  $C_s$ ; 5 CRs out  
304 of 16) or in the cuboid-smooth quadrants (low  $C_f$ , higher  $C_s$ ; 6 CRs out of 16). On the contrary, only  
305 few of the analysed products are found in the elongated-rough (high  $C_f$ , low  $C_s$ ; 3 out of 16) and  
306 elongated-smooth (high  $C_f$ , higher  $C_s$ ; 2 out of 16) quadrants.

307 It can be noticed that, as expected (Figure 2), cryogenic product G 0.2-0.6 is characterized by the  
308 highest value of  $C_s$ , typical of smooth particles obtained from the low-temperature and high-impact  
309 production procedure which causes brittle fracture of rubber granules. Results obtained for the CR  
310 derived from the waterjet process (H 0-0.7) are also coherent with physical expectations, with a  
311 similar shape ( $C_f$  of the order of 1.65) but a rougher surface caused by the abrasive action of high-  
312 pressure jets.

313 Figure 2. Examples of post-processing images of two CR samples.

314 Figure 3. Shape-roughness mapping plot of CR samples.

315 Figure 4. Effect of CR coarseness/fineness on surface area per unit mass.

316 Variability of  $C_f$ - $C_s$  results obtained for ambient size CRs is a consequence of the specific features of  
317 the different production processes. However, it can be observed that the coarser products, even

318 though regular in shape, tend to have a smoother surface ( $C_s > 0.85$ ). On the contrary, finer CR  
 319 products exhibit a rougher surface and the lowest  $C_s$  values are in fact associated to the lowest values  
 320 of parameter  $D_{50}$  (I 0.2-0.7, A 0.1-0.3, C 0.1-0.4, K 0.1-0.5 and B 0.1-0.4, all with  $D_{50} < 0.50$  mm).  
 321 The above described variability of particle size distribution and morphological characteristics has an  
 322 effect on the specific surface area (per unit mass) of analyzed CRs, which is reported in Tables 3a  
 323 and 3b.  
 324 As expected, the highest and lowest surface area values are respectively associated to the “fine” and  
 325 “coarse” products (with average values equal to  $0.0243 \text{ m}^2/\text{g}$  and  $0.0046 \text{ m}^2/\text{g}$ ), while “standard”  
 326 CRs exhibit intermediate  $SA_m$  values (mean equal to  $0.0127 \text{ m}^2/\text{g}$ ). Such particle size dependency is  
 327 visually represented in Figure 4, where surface area is plotted as a function of the diameter  
 328 corresponding to 50% passing ( $D_{50}$ ).  
 329 Results obtained from chemical analyses highlighted several common factors for all CRs, while  
 330 singularities were related to specific aspects of production, such as treatment (number and type of  
 331 ELT processing phases) and feed (type and conditions of ELTs, presence of non-ELT rubber).  
 332 Elemental analysis showed, as expected, that percentages of carbon and hydrogen are similar for all  
 333 CRs, with the exception of “coarse” ones (I 0.6-1.5 and J 0.9-2.2). Such a result is coherent with  
 334 density (and specific gravity) data, listed in Table 2, which exhibit a similar uniformity, with the  
 335 exception of cryogenic product G 0.2-0.6, and do not reveal an explicit dependency from the specific  
 336 features of production processes.  
 337 Nitrogen and sulphur contents, lower in absolute value, were found to be slightly more variable. The  
 338 latter element, used for rubber vulcanization, may be an indicator of the possible use of different  
 339 percentages of car vs. truck ELTs, of the selection of ELTs in specific conditions (depending upon  
 340 tread wear) and/or of the introduction in the processing plant of additional materials such as non-  
 341 vulcanized (raw) rubber and scrap rubber from other production processes.  
 342 Regardless of the elemental composition uniformity described above, CR samples were found to be  
 343 significantly different in terms of their level of cleanliness and purity, which was assessed by

344 evaluating content of metals by means of ICP-OES analyses. This can be appreciated by considering  
345 the experimental data provided in Table 5 and in Figures 5a and 5b, where only the most relevant  
346 elements have been displayed.

347 Whenever two different CRs were sampled from a same plant, metal content was found to be higher  
348 in the finer fraction (i.e. products from plants A, B, C, I and J). This is due to the fact that as particle  
349 size is reduced, separation of different components (rubber, steel, textiles) is more difficult.

350 Figure 5a. Metal content of CR samples.

351 Figure 5b. Metal content of CR samples.

352 By looking at the entire data set it can also be observed that the lowest metals' content is exhibited  
353 by CRs J 0.9-2.2 ("coarse") and F 0.3-0.7 ("standard"), while the highest values were found in the  
354 case of C 0.1-0.4 ("fine"). For such products overall ranking deriving from considering all major  
355 elements was matched by the assessment focused on iron content, which ranged from 0.036-0.042%  
356 (CRs J 0.9-2.2 and F 0.3-0.7) to 0.915% (CR C 0.1-0.4).

357 Singularities were detected for CRs produced by plants D and G, which were characterized by very  
358 high values of lead (D 0.3-0.7) and calcium (G 0.2-0.6) content. Moreover, both products were found  
359 to have out-of-range contents of barium. Anomalies of the barium content may be related to its  
360 employment as a catalyst for the synthesis of polybutadiene rubber, while presence of excessive  
361 calcium may derive from calcium hydroxide used for the anti-packing treatment of non-vulcanized  
362 rubber which may have been introduced in the production flow of CR. Finally, probably as a result  
363 of inefficient separation of residues coming from steel belts and bead wires, an exceptionally high  
364 copper content was detected in CR C 0.1-0.4.

365 All products were found to have a zinc content greater than 1%, presumably due to the  
366 corresponding oxide that is used as a vulcanisation aid in the rubber production process.

367 The significant variability of plant configuration in the case of ambient size reduction processes does  
368 not allow a straightforward comparison between the three technologies in terms of cleanliness and



369 purity. However, results obtained for products G 0.2-0.6 and H 0-0.7 suggest that the cryogenic  
370 treatment may provide a better means for rubber-metal separation than waterjet blasting.

371 The observations provided above, based on analytical measurements, were supplemented by visual  
372 analyses of microscopic images which focused on the detection of foreign impurities (metals and  
373 textiles) not included in the rubber matrix. Examples of the assessment performed on the two CRs  
374 produced by plant C are given in Figure 6.

375 CR samples were subjected to laboratory tests for the determination of VOCs and PAHs since there  
376 are concerns whether the presence of recycled rubber in bituminous mixtures can affect the toxic and  
377 carcinogenic potential of gaseous emissions produced during laying on site. However, it should be  
378 pointed out that such an analysis was included in this study for comparative purposes only, since the  
379 Authors have already highlighted the fact that the dominating factors which control VOCs and PAHs  
380 released in the environment are bitumen type and quantity (Zanetti et al., 2013a and 2014a).

381 Results of VOC and PAH analyses are listed in Tables 6 and 7 and are synthetically represented, by  
382 referring to total contents, in Figure 7.

383 Figure 6. Examples of visual assessment of degree of cleanliness of two CR samples.

384 In the case of ambient size CRs, it can be postulated that results are dependent not only on plant feed  
385 (type and wear conditions of ELTs, possible presence of non-ELT rubber) but also on two additional  
386 factors which have a direct effect on the release of organic compounds in the environment during CR  
387 production. Both factors should be taken into account since they may cause a reduction of VOCs and  
388 PAHs which can be detected in the laboratory on the final product.

389 The first relevant factor is energy dissipation, which occurs, in the form of heat, during shredding  
390 and milling operations. Thus, as the number of such phases is increased, for example when passing  
391 from “coarse” to “standard” (or “fine”) products, lighter organic compounds in the CR are bound to  
392 be reduced. This is the case of products derived from plants I and J, which exhibit a total VOC  
393 content which decreases when considering finer products obtained by supplementary milling.

394 However, it is uncertain whether the same explanation can be given for VOC data obtained on

products coming from plants A, B and C, since the fine fractions may have been derived from the “standard” ones only by means of additional sieving (which causes no heating effects). The second relevant factor is specific surface area. In fact, for a given production temperature, loss of VOCs is presumably more intense as specific surface increases and, as a consequence, CRs with a higher  $SA_m$  are expected to yield lower VOCs in laboratory analyses. As shown in Figure 8, such a physical expectation is matched by results found for CRs from plants A, B and C (and, to a more limited extent, from I and J) being the finer products those which possess a higher  $SA_m$  value.

Figure 7. VOC and PAH content of CR samples.

Figure 8. Effect of surface area on total VOC content of CR samples.

PAH contents do not show the same straightforward particle size and surface area dependencies of VOCs. This is coherent with the fact that such compounds, in the typical conditions which occur during ELT processing, are less prone to be released in the environment and are therefore mostly maintained within the rubber matrix of the final product.

Results obtained on CRs derived from cryogenic and waterjet production (G 0.2-0.6 and H 0-0.7) cannot be explained by referring to the same hypotheses outlined above since milling occurs in temperature conditions which are more tightly controlled. Thus, they should be considered mainly as a consequence of the characteristics of materials fed to production and do not allow any interpretation theory to be proposed.

## 5. CONCLUSIONS

Experimental data collected in the study described in this paper show that CRs for paving applications may have, depending upon their production process, physical and chemical properties which are comprised within very broad variation ranges. Such an observation is extremely relevant since significant variations may be reflected, as a consequence, in the bituminous materials in which CRs are included (either “wet” asphalt rubber binders or “dry” bituminous mixtures). Thus, several

420 key elements should be taken into account in the production of CRs and in their performance-  
421 oriented selection.

422 In order to maximize bitumen-CR interactions, which are of premium importance in asphalt rubber  
423 binders, evaluation of specific surface area should be included in CR standard characterization  
424 procedures. In fact, experimental data show that surface area depends upon particle size distribution,  
425 which can be controlled by adequately combining shredding, milling and sieving operations, and by  
426 particle morphology, which can be mapped by referring to image analysis parameters.

427 Cleanliness of CR products also plays a critical role in the performance of final binders and mixtures  
428 since textile and metal residues may impair homogeneity and act as damage initiators. In such a  
429 context, combined analysis of the metals' content and results of microscopic observations may be of  
430 support in identifying specific foreign elements and oxides and in implementing improvement  
431 measures in the processing of ELTs.

432 Chemical characterization of CRs with respect to organic compounds such as VOCs and PAHs is  
433 also of special interest as a result of their possible effects on gaseous emissions produced by  
434 composite bituminous binders and mixtures during laying operations. Although it has been already  
435 proven that emissions are more strongly affected by bitumen type and content and that the potential  
436 release of CR is in fact shielded by bitumen itself, analyses performed on CR may be of value at  
437 least in comparative terms. As highlighted in this paper, a reduction of such a potential may be  
438 obtained by selecting products derived from a greater number of shredding/milling phases and/or  
439 characterized by a higher specific surface area.

440 Further investigations are currently being performed in order to widen the existing CR database and  
441 to highlight in more detail relationships between CR characteristics and production features. A CR  
442 certification process is also under development with the purpose of providing the paving sector with  
443 an efficient tool for the selection of materials, processes and products.

444

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449

Table 2. Size distribution parameters, density and specific gravity of the CR samples.

|   | A       | A       | B       | B       | C       | C       | D       | E       | F       | G       | H     | I       | I       | J       | J       | K       |
|---|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|-------|---------|---------|---------|---------|---------|
|   | 0.4-0.7 | 0.1-0.3 | 0.3-0.7 | 0.1-0.4 | 0.4-0.7 | 0.1-0.4 | 0.3-0.7 | 0.3-0.6 | 0.3-0.7 | 0.2-0.6 | 0-0.7 | 0.6-1.5 | 0.2-0.7 | 0.9-2.2 | 0.3-0.6 | 0.1-0.5 |
| Diameter with 10% passing ( $D_{10}$ ) [mm]     | 0.42    | 0.11    | 0.30    | 0.15    | 0.38    | 0.09    | 0.27    | 0.27    | 0.27    | 0.24    | 0     | 0.57    | 0.17    | 0.94    | 0.25    | 0.13    |
| Diameter with 90% passing ( $D_{90}$ ) [mm]     | 0.69    | 0.33    | 0.68    | 0.38    | 0.69    | 0.39    | 0.70    | 0.55    | 0.67    | 0.56    | 0.74  | 1.50    | 0.69    | 2.21    | 0.56    | 0.54    |
| Diameter with 50% passing ( $D_{50}$ ) [mm]     | 0.54    | 0.23    | 0.51    | 0.27    | 0.54    | 0.23    | 0.51    | 0.45    | 0.50    | 0.42    | 0.28  | 1.04    | 0.41    | 1.48    | 0.35    | 0.29    |
| Uniformity coefficient $D_{80}/D_{20}$          | 1.41    | 1.86    | 1.65    | 1.83    | 1.44    | 2.50    | 1.60    | 1.58    | 1.62    | 1.80    | 8.03  | 1.89    | 2.48    | 1.78    | 1.82    | 2.60    |
| Percentage passing 0.063 mm ( $P_{0.063}$ ) [%] | 0.07    | 6.47    | 0.58    | 59.9    | 0.33    | 34.8    | 0.97    | 0.61    | 0.13    | 21.0    | 14.3  | 0       | 0.75    | 0       | 0       | 2.31    |
| Density ( ) [g/cm <sup>3</sup> ]                | 1.172   | 1.213   | 1.181   | 1.192   | 1.158   | 1.196   | 1.203   | 1.178   | 1.185   | 1.223   | 1.189 | 1.204   | 1.199   | 1.207   | 1.190   | 1.208   |
| Specific gravity (SG)                           | 1.174   | 1.215   | 1.183   | 1.194   | 1.160   | 1.200   | 1.206   | 1.181   | 1.188   | 1.225   | 1.192 | 1.207   | 1.201   | 1.210   | 1.193   | 1.210   |

Table 3a. Morphological parameters and surface area of the CR samples (plants A to E).

|   | A       | A       | B       | B       | C       | C       | D       | E       |
|---|---------|---------|---------|---------|---------|---------|---------|---------|
|   | 0.4-0.7 | 0.1-0.3 | 0.3-0.7 | 0.1-0.4 | 0.4-0.7 | 0.1-0.4 | 0.3-0.7 | 0.3-0.6 |
| Shape coefficient ( $C_f$ )                               | 1.79    | 1.77    | 1.85    | 1.69    | 1.69    | 1.78    | 1.67    | 1.52    |
| Solidity coefficient ( $C_s$ )                            | 0.848   | 0.812   | 0.851   | 0.834   | 0.855   | 0.805   | 0.846   | 0.871   |
| Shape corrective factor ( $\phi_f$ )                      | 0.955   | 0.973   | 0.918   | 0.924   | 0.978   | 0.930   | 0.833   | 0.894   |
| Roughness factor ( $\phi_r$ )                             | 1.240   | 1.313   | 1.216   | 1.206   | 1.254   | 1.244   | 1.078   | 1.091   |
| Corrective factor ( $\phi$ )                              | 1.184   | 1.277   | 1.117   | 1.115   | 1.226   | 1.157   | 0.897   | 0.975   |
| Surface area per unit mass ( $SA_m$ ) [m <sup>2</sup> /g] | 0.0112  | 0.0270  | 0.0111  | 0.0208  | 0.0118  | 0.0252  | 0.0088  | 0.0110  |

Table 3b. Morphological parameters and surface area of the CR samples (plants F to L).

|   | F       | G       | H      | I       | I       | J       | J       | K       |
|---|---------|---------|--------|---------|---------|---------|---------|---------|
|   | 0.3-0.7 | 0.2-0.6 | 0-0.7  | 0.6-1.5 | 0.2-0.7 | 0.9-2.2 | 0.3-0.6 | 0.1-0.5 |
| Shape coefficient ( $C_f$ )                       | 1.63    | 1.63    | 1.66   | 1.65    | 1.74    | 1.54    | 1.76    | 1.63    |
| Solidity coefficient ( $C_s$ )                    | 0.879   | 0.899   | 0.809  | 0.868   | 0.812   | 0.885   | 0.871   | 0.819   |
| Shape corrective factor ( $\phi_f$ )              | 0.927   | 0.878   | 0.745  | 0.933   | 0.967   | 0.950   | 0.919   | 0.950   |
| Roughness factor ( $\phi_r$ )                     | 1.166   | 1.078   | 1.017  | 1.201   | 1.315   | 1.228   | 1.158   | 1.311   |
| Corrective factor ( $\phi$ )                      | 1.080   | 0.946   | 0.758  | 1.120   | 1.271   | 1.167   | 1.064   | 1.245   |
| Surface area per unit mass ( $SA_m$ ) [ $m^2/g$ ] | 0.0109  | 0.0111  | 0.0136 | 0.0054  | 0.0151  | 0.0039  | 0.0149  | 0.0205  |

Table 4. C, H, N and S contents of the CR samples.

|       | A       | A       | B       | B       | C       | C       | D       | E       | F       | G       | H     | I       | I       | J       | J       | K       |
|-------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|-------|---------|---------|---------|---------|---------|
|       | 0.4-0.7 | 0.1-0.3 | 0.3-0.7 | 0.1-0.4 | 0.4-0.7 | 0.1-0.4 | 0.3-0.7 | 0.3-0.6 | 0.3-0.7 | 0.2-0.6 | 0-0.7 | 0.6-1.5 | 0.2-0.7 | 0.9-2.2 | 0.3-0.6 | 0.1-0.5 |
| C (%) | 77.00   | 77.10   | 79.22   | 78.83   | 78.26   | 76.21   | 81.89   | 81.54   | 78.37   | 77.05   | 81.50 | 82.10   | 81.50   | 84.40   | 80.80   | 81.60   |
| H (%) | 7.24    | 7.02    | 7.34    | 7.16    | 7.38    | 7.00    | 7.23    | 7.25    | 7.03    | 7.09    | 7.42  | 7.41    | 7.35    | 7.55    | 7.27    | 7.61    |
| N (%) | 0.52    | 0.51    | 0.47    | 0.48    | 0.49    | 0.53    | 0.48    | 0.39    | 0.46    | 0.43    | 0.43  | 0.42    | 0.54    | 0.37    | 0.46    | 0.43    |
| S (%) | 2.14    | 1.86    | 2.02    | 1.98    | 2.42    | 2.22    | 2.33    | 1.69    | 2.03    | 1.96    | 1.86  | 1.80    | 1.95    | 1.84    | 2.14    | 1.48    |

Table 5. Metal contents of the CR samples.

|    |         | A       | A       | B       | B       | C       | C       | D       | E       | F       | G       | H     | I       | I       | J       | J       | K       |
|----|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|-------|---------|---------|---------|---------|---------|
|    |         | 0.4-0.7 | 0.1-0.3 | 0.3-0.7 | 0.1-0.4 | 0.4-0.7 | 0.1-0.4 | 0.3-0.7 | 0.3-0.6 | 0.3-0.7 | 0.2-0.6 | 0-0.7 | 0.6-1.5 | 0.2-0.7 | 0.9-2.2 | 0.3-0.6 | 0.1-0.5 |
| Na | (mg/kg) | 216     | 326     | 240     | 231     | 214     | 218     | 230     | 198     | 229     | 252     | 283   | 219     | 317     | 202     | 203     | 251     |
| K  | (mg/kg) | 506     | 809     | 504     | 586     | 530     | 582     | 559     | 407     | 300     | 357     | 1514  | 738     | 1105    | 628     | 745     | 1030    |
| Ca | (%)     | 0.349   | 0.411   | 0.364   | 0.546   | 0.180   | 0.357   | 0.345   | 0.496   | 0.130   | 1.120   | 0.181 | 0.268   | 0.276   | 0.235   | 0.382   | 0.354   |
| Mg | (mg/kg) | 444     | 719     | 350     | 542     | 397     | 999     | 445     | 1240    | 246     | 344     | 390   | 352     | 526     | 281     | 379     | 569     |
| Fe | (%)     | 0.153   | 0.440   | 0.088   | 0.245   | 0.169   | 0.915   | 0.215   | 0.223   | 0.042   | 0.147   | 0.320 | 0.549   | 0.388   | 0.036   | 0.172   | 0.125   |
| Mn | (mg/kg) | 14.7    | 34.9    | 10.7    | 23.9    | 16.1    | 59.6    | 19.6    | 25.3    | 5.1     | 15.8    | 26.4  | 34.0    | 29.9    | 4.7     | 13.5    | 14.5    |
| Ba | (mg/kg) | 13.2    | 18.5    | 8.87    | 20.3    | 10.9    | 28.3    | 211.0   | 18.7    | 6.3     | 121.0   | 12.5  | 10.6    | 10.9    | 8.2     | 10.7    | 11.3    |
| Al | (mg/kg) | 630     | 1047    | 572     | 779     | 493     | 707     | 800     | 675     | 372     | 653     | 572   | 534     | 908     | 463     | 690     | 641     |
| Cd | (mg/kg) | 4.59    | 4.11    | 6.30    | 4.48    | 5.79    | 5.64    | 3.40    | 2.89    | 2.17    | 2.43    | 3.04  | 5.64    | 4.26    | 5.93    | 4.76    | 5.11    |
| Cr | (mg/kg) | 4.73    | 8.65    | 7.02    | 6.27    | 6.69    | 11.7    | 5.07    | 12.3    | 2.29    | 3.51    | 46.4  | 5.00    | 27.2    | 2.8     | 4.3     | 5.6     |
| Ni | (mg/kg) | 11.5    | 8.12    | 10.2    | 9.13    | 9.87    | 13.2    | 9.22    | 11.0    | 3.84    | 4.54    | 28.3  | 8.54    | 13.1    | 4.7     | 7.2     | 9.4     |
| Pb | (mg/kg) | 66.3    | 32.8    | 77.7    | 44.9    | 73.3    | 73.5    | 194.0   | 26.6    | 28.4    | 28.0    | 22.7  | 40.0    | 35.2    | 28.6    | 40.0    | 24.0    |
| Cu | (mg/kg) | 295     | 656     | 222     | 472     | 317     | 1218    | 353     | 80.0    | 64.3    | 85.9    | 118   | 149     | 447     | 32      | 298     | 73      |
| Zn | (%)     | 2.03    | 1.21    | 1.94    | 1.83    | 2.10    | 2.26    | 1.87    | 1.33    | 1.16    | 1.18    | 1.41  | 1.50    | 1.35    | 1.34    | 1.54    | 1.25    |
| Co | (mg/kg) | 330     | 232     | 248     | 259     | 347     | 418     | 255     | 179     | 162     | 151     | 261   | 240     | 218     | 183     | 317     | 36.2    |
| Ti | (mg/kg) | 55.5    | 53.6    | 49.7    | 67.4    | 33.6    | 45.1    | 56.0    | 39.6    | 37.0    | 65.2    | 31.2  | 41.5    | 37.9    | 37.8    | 49.5    | 36.3    |
| Sb | (mg/kg) | 487     | 229     | 367     | 379     | 554     | 608     | 388     | 183     | 164     | 151     | 258   | 247     | 214     | 191     | 327     | 36.3    |

Table 6. VOC contents (in mg/kg) of the CR samples.

|                        | A       | A       | B       | B       | C       | C       | D       | E       | F       | G       | H      | I       | I       | J       | J       | K       |
|------------------------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|--------|---------|---------|---------|---------|---------|
|                        | 0.4-0.7 | 0.1-0.3 | 0.3-0.7 | 0.1-0.4 | 0.4-0.7 | 0.1-0.4 | 0.3-0.7 | 0.3-0.6 | 0.3-0.7 | 0.2-0.6 | 0-0.7  | 0.6-1.5 | 0.2-0.7 | 0.9-2.2 | 0.3-0.6 | 0.1-0.5 |
| benzene                | 17.24   | 7.56    | 33.68   | 15.63   | 29.41   | 15.18   | 10.10   | 18.43   | 16.32   | 8.94    | 4.81   | 51.08   | 14.47   | 34.03   | 31.30   | 13.29   |
| toluene                | 4.04    | 1.59    | 1.60    | 1.20    | 2.68    | 1.58    | 0.01    | 1.68    | 1.56    | 1.52    | 1.07   | 1.66    | 1.74    | 0.52    | 0.56    | 0.46    |
| ethylbenzene           | 1.35    | 1.44    | 1.78    | 1.10    | 1.15    | 2.06    | 1.35    | 2.45    | 3.70    | 0.59    | 1.80   | 1.40    | 1.36    | 1.08    | 1.33    | 2.26    |
| styrene                | 0.34    | 0.23    | 0.14    | 0.11    | 0.24    | 0.52    | 0.05    | 0.40    | 0.18    | 0.11    | 0.16   | 0.23    | 0.14    | 0.06    | 0.06    | 0.03    |
| 1,2,4-trimethylbenzene | 4.77    | 0.82    | 2.84    | 2.44    | 1.60    | 1.65    | 1.80    | 4.08    | 2.78    | 2.84    | 1.24   | 5.50    | 0.88    | 0.94    | 0.97    | 0.92    |
| 1,3,5-trimethylbenzene | 1.35    | 1.46    | 0.95    | 1.32    | 0.61    | 1.10    | 1.85    | 1.58    | 1.68    | 1.66    | 1.48   | 3.01    | 1.74    | 1.04    | 1.21    | 0.42    |
| p-xylene               | 1.88    | 1.09    | 1.55    | 1.04    | 1.52    | 1.46    | 1.24    | 0.92    | 1.52    | 1.16    | 1.67   | 2.10    | 1.22    | 1.34    | 1.57    | 0.24    |
| 1,3,5-trichlorobenzene | 5.84    | 1.25    | 3.02    | 3.64    | 3.20    | 1.19    | 4.43    | 1.56    | 4.53    | 3.44    | 1.14   | 5.61    | 2.30    | 0.94    | 0.97    | 0.92    |
| 1,2,4-trichlorobenzene | 0.34    | <0.001  | <0.001  | <0.001  | 0.33    | <0.001  | 0.31    | 0.62    | 0.34    | 0.58    | <0.001 | 0.760   | <0.001  | 0.180   | 0.140   | 0.230   |
| Total VOCs             | 37.15   | 15.44   | 45.56   | 26.48   | 40.74   | 24.74   | 21.14   | 31.72   | 32.61   | 20.84   | 13.37  | 71.35   | 23.85   | 40.13   | 38.11   | 18.77   |



Table 7. PAH contents (in mg/kg) of the CR samples.

|                         | A       | A       | B       | B       | C       | C       | D       | E       | F       | G       | H      | I       | I       | J       | J       | K       |
|-------------------------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|--------|---------|---------|---------|---------|---------|
|                         | 0.4-0.7 | 0.1-0.3 | 0.3-0.7 | 0.1-0.4 | 0.4-0.7 | 0.1-0.4 | 0.3-0.7 | 0.3-0.6 | 0.3-0.7 | 0.2-0.6 | 0-0.7  | 0.6-1.5 | 0.2-0.7 | 0.9-2.2 | 0.3-0.6 | 0.1-0.5 |
|                         | 0.30    | 0.22    | 0.27    | 0.21    | 0.42    | 0.34    | 0.34    | 0.53    | 0.32    | 0.37    | 0.49   | 0.40    | 0.32    | 0.83    | 0.71    | 1.06    |
| naphtalene              |         |         |         |         |         |         |         |         |         |         |        |         |         |         |         |         |
| acenaphtylene           | 0.32    | 0.44    | 0.35    | 0.20    | 0.61    | 0.51    | 0.45    | 0.74    | 0.46    | 0.47    | 0.72   | 0.55    | 0.68    | 0.54    | 0.58    | 0.64    |
| 1-bromonaphtalene       | 0.03    | 0.02    | 0.05    | 0.05    | 0.07    | 0.06    | 0.06    | 0.02    | 0.06    | 0.05    | 0.02   | 0.16    | 0.01    | 0.12    | 0.07    | 0.13    |
| acenaphtene             | 0.06    | 0.11    | 0.06    | 0.04    | 0.08    | 0.06    | 0.07    | 0.08    | 0.06    | 0.06    | 0.06   | 0.13    | 0.08    | 0.04    | 0.04    | 0.04    |
| fluorene                | 0.02    | 0.28    | 0.16    | 0.12    | 0.22    | 0.21    | 0.20    | 0.28    | 0.19    | 0.21    | 0.12   | 0.35    | 0.02    | 0.21    | 0.17    | 0.03    |
| phenantrene             | 2.80    | 2.84    | 1.45    | 1.56    | 1.90    | 1.84    | 1.68    | 3.36    | 1.54    | 2.27    | 1.88   | 3.38    | 1.80    | 4.15    | 4.36    | 1.20    |
| anthracene              | 1.80    | 0.53    | 0.47    | 0.32    | 0.79    | 0.38    | 0.60    | 0.91    | 0.61    | 0.87    | 0.54   | 5.03    | 3.62    | 6.49    | 7.05    | 5.40    |
| fluorantene             | 4.08    | 3.93    | 2.72    | 3.02    | 3.14    | 2.36    | 3.30    | 3.13    | 2.96    | 3.38    | 3.59   | 0.91    | 0.98    | 0.31    | 0.27    | 0.15    |
| pyrene                  | 13.12   | 12.51   | 10.02   | 10.52   | 11.30   | 17.20   | 12.46   | 16.04   | 11.29   | 10.92   | 10.46  | 17.90   | 13.30   | 18.43   | 24.73   | 9.09    |
| benzo[a]anthracene      | 2.34    | 1.74    | 1.62    | 1.52    | 0.87    | 0.80    | 2.34    | 1.13    | 1.77    | 3.02    | 0.68   | 1.40    | 1.88    | 0.75    | 0.61    | 0.12    |
| triphenylene            | 0.51    | 0.42    | 0.22    | 0.30    | 0.12    | 0.12    | 0.34    | 0.16    | 0.23    | 0.88    | 0.12   | 0.29    | 0.38    | 0.85    | 0.80    | 0.33    |
| benzo[a]pyrene          | 0.85    | 0.60    | 0.85    | 0.45    | 0.47    | 1.22    | 1.22    | 0.61    | 1.02    | 1.85    | 0.45   | 0.41    | 1.06    | 1.91    | 2.67    | 0.53    |
| benzo[b]fluorantene     | 0.94    | 0.58    | 0.67    | 0.48    | 0.31    | 1.21    | 0.96    | 1.43    | 0.76    | 1.56    | 0.31   | 0.59    | 0.76    | 1.67    | 2.83    | 1.08    |
| dibenzo [a,h]anthracene | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  | <0.001 | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  |
| indeno[1,2,3-cd]pyrene  | 0.33    | <0.001  | 0.19    | 0.15    | 0.10    | 0.20    | 0.12    | 0.14    | 0.10    | 0.19    | <0.001 | <0.001  | <0.001  | <0.001  | <0.001  | <0.001  |
| benzo[ghi]perylene      | 3.68    | 1.52    | 3.11    | 1.82    | 1.35    | 0.84    | 3.62    | 2.04    | 3.54    | 4.56    | 0.80   | 2.63    | 2.60    | 2.99    | 3.25    | 0.34    |
| Total PAHs              | 31.18   | 25.74   | 22.21   | 20.76   | 21.75   | 27.35   | 27.76   | 30.60   | 24.91   | 30.66   | 20.24  | 34.13   | 27.49   | 39.29   | 48.14   | 20.14   |

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